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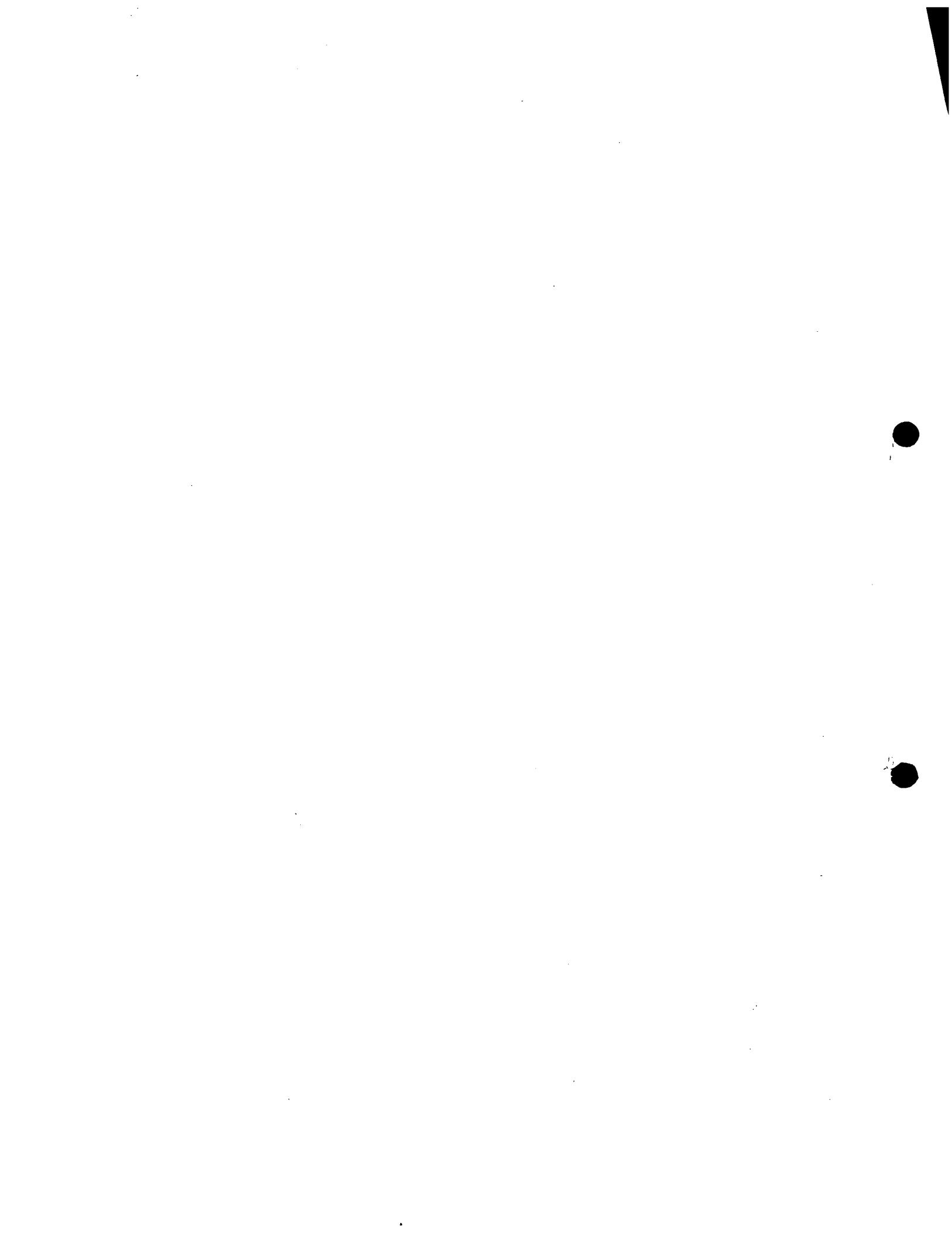
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Signed *Anne Bowen*

Dated 2 November 1999



Patents Act 1977
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18 DEC 1998

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

The Patent Office

Cardiff Road
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Gwent NP9 1RH

1. Your Reference

MG/AJT/PG3612

2. Patent application number

(The Patent office will fill in this part)

9828074.6

3. Full name, address and postcode of the or of each applicant (underline all surnames)

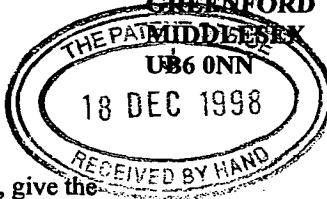
GLAXO GROUP LIMITED
GLAXO WELLCOME HOUSE
BERKELEY AVENUE

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18 DEC 1998



Patents ADP number (if you know it)

21 DEC 98 E413325-1 D01030
P01/7700 0.00 - 9828074.6

If the applicant is a corporate body, give the country/state of its corporation

THERAPEUTICALLY USEFUL COMPOUNDS

5 Name of your agent (if you know one)

ANDREW J. TEUTEN
(SEE CONTINUATION SHEET)

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GLAXO WELLCOME PLC
GLAXO WELLCOME HOUSE, BERKELEY AVENUE
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UB6 0NN, GB

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7027683 001

6. If you are declaring priority from one or more earlier patent applications, give the country and date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)Date of Filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)8. Is a statement of inventorship and of right to grant a patent required in support of this request? (Answer yes if:
a) any applicant named in part 3 is not an inventor, or
b) there is an inventor who is not named as an applicant, or
c) any named applicant is a corporate body.

See note (d)

9. Enter the number of sheets for any of the following items you are filing with this form.
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Continuation sheets of this form	1
Description	92
Claim(s)	5
Abstract	-
Drawing(s)	-

10. If you are also filing any of the following, state how many against each item

Priority Documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (*Patents Form 7/77*)

Request for preliminary examination and search (*Patent Form 9/77*)

Request for substantive examination (*Patent Form 10/77*)

Any other documents
(please specify)

11.

I/We request the grant of a patent on the basis of this application

Signature: Andrew J. Teuten
AGENT FOR THE APPLICANTS

18 December 1998

12. Name and daytime telephone number of person to contact in the United Kingdom

Fiona Longstaff
0181-966 8234

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Therapeutically useful compounds

This invention relates to novel chemical compounds, processes for their preparation, pharmaceutical formulations containing them and their use in therapy.

Inflammation is a primary response to tissue injury or microbial invasion and is characterised by leukocyte adhesion to the endothelium, diapedesis and activation within the tissue. Leukocyte activation can result in the generation of toxic oxygen species (such as superoxide anion), and the release of granule products (such as peroxidases and proteases). Circulating leukocytes include neutrophils, eosinophils, basophils, monocytes and lymphocytes. Different forms of inflammation involve different types of infiltrating leukocytes, the particular profile being regulated by the profile of adhesion molecule, cytokine and chemotactic factor expression within the tissue.

The primary function of leukocytes is to defend the host from invading organisms, such as bacteria and parasites. Once a tissue is injured or infected, a series of events occurs which causes the local recruitment of leukocytes from the circulation into the affected tissue. Leukocyte recruitment is controlled to allow for the orderly destruction and phagocytosis of foreign or dead cells, followed by tissue repair and resolution of the inflammatory infiltrate. However in chronic inflammatory states, recruitment is often inappropriate, resolution is not adequately controlled and the inflammatory reaction causes tissue destruction.

Integrins are cell surface heterodimeric proteins comprising α and β chains, involved in the inflammatory process. The $\alpha 4$ -integrins, which include $\alpha 4\beta 1$ (also known as very late antigen-4 (VLA-4) or CD49d/CD29) and $\alpha 4\beta 7$, are expressed

mainly on leukocytes other than neutrophils (eg. eosinophils, T- and B-lymphocytes, basophils and mast cells). The adhesion molecule ligands for $\alpha 4$ -integrins include (i) the vascular cell adhesion molecule (VCAM-1; CD106), (ii) a sequence within the alternatively spliced connecting segment-1 (CS-1) in fibronectin (an extracellular matrix protein), and (iii) a site on the mucosal addressin cell adhesion molecule (MAdCAM). Under normal conditions, VCAM-1 is minimally expressed in the vasculature, however, upregulation of VCAM-1 on endothelial cells occurs near sites of inflammation. VCAM-1 has also been identified on a range of non-vascular cells including dendritic cells, bone marrow stromal cells, synoviocytes, astrocytes and some cortical neurons. MAdCAM expression is predominantly associated with gut tissue being expressed in the high endothelial veins of gut associated lymphoid tissue, peripheral lymph nodes and Peyers Patches.

Both $\alpha 4\beta 1$ (VLA-4) and $\alpha 4\beta 7$ can interact with VCAM-1, CS-1 in fibronectin and MAdCAM. The $\alpha 4$ -integrin/VCAM-1 interaction enables adhesion and subsequent transmigration of leukocytes through the wall of post-capillary venules to sites of tissue inflammation. Such an interaction is similarly capable of providing a co-stimulatory signal for T-cell activation, whilst the $\alpha 4$ -integrin/fibronectin interaction is believed to have a stimulatory role in the degranulation of mast cells, basophils and eosinophils. Therefore, $\alpha 4$ -integrin antagonists are capable of intervention at two levels to effect attenuation of the inflammatory processes which are essential in the pathophysiology of many chronic diseases. These include (i) inhibition of the recruitment of leukocytes to sites of tissue inflammation and (ii) inhibition of the activation of leukocytes and the release of inflammatory mediators.

Cell adhesion and signalling, mediated by $\alpha 4$ -integrins, are essential in numerous physiological and pathophysiological processes. The therapeutic

potential of $\alpha 4$ -integrin blocking agents has been investigated previously by testing specific $\alpha 4$ -integrin blocking monoclonal antibodies (anti- $\alpha 4$ -mAbs) in experimental *in vitro* and *in vivo* models of disease (Lobb and Hemler, 1994).

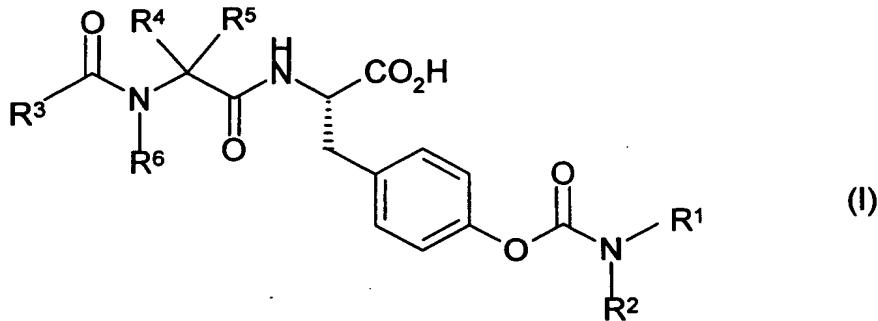
5 Anti- $\alpha 4$ -mAbs have shown beneficial effects in animal models of allergic lung inflammation relevant to asthma, including guinea-pig, rat, rabbit and sheep models. Additionally, anti- $\alpha 4$ -mAbs have also been shown to be efficacious in (i) rat and mouse models of experimental allergic encephalomyelitis (considered to be a model of the T-cell dependent autoimmune disease, multiple sclerosis), (ii) mouse models of contact hypersensitivity, (iii) colitis in the Cotton-top tamarin, 10 relevant to inflammatory bowel disease (Podolsky *et al*, 1993), and (iv) insulin dependent diabetes mellitus in the non-obese diabetic mouse (Baron *et al*, 1994). Fibronectin-derived peptides which are thought to block $\alpha 4$ -integrin function have shown efficacy in mouse contact hypersensitivity (Ferguson *et al*, 1991) and in rat adjuvant arthritis (Wahl *et al*, 1994).

15

We have now found a novel group of $\alpha 4$ -integrin antagonist compounds, with the potential to block leukocyte adhesion and activation, consequently affecting anti-inflammatory properties. These compounds are therefore of potential therapeutic benefit, especially in providing protection from leukocyte-induced 20 tissue damage in diseases where leukocytes are implicated at the site of inflammation.

25

Thus, according to one aspect of the invention, we provide compounds of formula I:



5

wherein R¹ and R² independently represent

(i) -C₁₋₆ alkyl, -C₃₋₈ cycloalkyl or -C₁₋₃ alkylC₃₋₈ cycloalkyl,

or such a group in which alkyl or cycloalkyl is substituted by one or more halogen, -CN, nitro, hydroxy or -OC₁₋₆alkyl groups;

(ii) -(CH₂)_eAr¹ or -(CH₂)_eOAr¹;
or NR¹R² together represent pyrrolidinyl, piperidinyl, piperazinyl, thiomorpholinyl, morpholinyl or azepinyl, or such a group fused to a benzene ring, optionally substituted by one or more -(CO)_n(CH₂)_tAr¹, -(CO)_nC₁₋₆ alkylAr¹Ar², -(CO)_nC₁₋₆ alkyl, -(CH₂)_rOH, -(CH₂)_rO(CH₂)_pOH, -(CH₂)_rOC₁₋₆ alkyl, -O(CH₂)_tAr¹, -(CH₂)_rSO₂Ar¹, piperidin-1-yl, -(CH₂)_tCONR⁸R⁹, -NR¹⁰(CO)_n(CH₂)_tAr¹, -NR¹⁰(CO)_nC₁₋₃ alkylC₃₋₆ cycloalkyl, -NR¹⁰(CO)_nC₁₋₆ alkylC₃₋₆ cycloalkyl, -CONR¹⁰(CH₂)_tAr¹, halogen, -NHSO₂C₁₋₆ alkyl, -SO₂NR¹⁰R¹¹, -SO₂C₁₋₆ alkyl or -SO₂Ar² groups;

20 R³ represents -C₁₋₆alkylNHC(=NH)NH₂, -C₂₋₆alkenylNHC(=NH)NH₂, -C₂₋₆alkynylNHC(=NH)NH₂, -C₁₋₆alkylNR¹⁴R¹⁸, -(CH₂)_hCONR¹⁴R¹⁸, -(CH₂)_hCOC₁₋₆ alkyl, -(CH₂)_dCHNR¹⁸CONR²⁰R²¹, -(CH₂)_mNR¹⁸CONR¹⁴R¹⁸, -(CH₂)_dNR¹⁸Ar³, -(CH₂)_dCONR¹⁸Ar³, -(CH₂)_hCOOR¹⁸, -(CH₂)_cAr³, -O(CH₂)_cAr³, -(CH₂)_dCO(CH₂)_sAr³ or -(CH₂)_dOAr³;

or R³ represents -(CH₂)_c-2,4-imidazolidinedione, -(CH₂)_c(piperidin-4-yl), -(CH₂)_c(piperidin-3-yl), -(CH₂)_c(piperidin-2-yl), -(CH₂)_c(morpholin-3-yl) or -(CH₂)_c(morpholin-2-yl) optionally substituted on nitrogen by -(CO)_fC₁₋₆alkyl, -(CO)_f(CH₂)_cAr² or -C(=NH)NH₂;

5 or R³ represents -(CH₂)_ddibenzofuran optionally substituted by -C₁₋₆alkyl or halogen;

or R³ represents -(CH₂)_c-thioxanthen-9-one;

R⁴ represents hydrogen, -C₁₋₆ alkyl, -C₁₋₃ alkylC₃₋₆ cycloalkyl, -(CH₂)_qAr², -C₁₋₄alkyl-X-R⁷, -C₁₋₄alkyl SO₂C₁₋₄ alkyl, -C₁₋₆alkylNR¹²R¹³ or -C₁₋₆ alkylNR¹²COC₁₋₆ alkyl;

10 R⁵ represents hydrogen, or R⁴R⁵ together with the carbon to which they are attached form a C₅₋₇ cycloalkyl ring;

15 R⁶ represents hydrogen or -C₁₋₆alkyl, or R⁶ and R⁴ together with the N and C atoms to which they are respectively attached form a pyrrolidine ring;

20 R⁷ represents hydrogen, -(CH₂)_wNR¹²R¹³, -(CH₂)_uAr² or -(CH₂)_wNR¹²COC₁₋₆ alkyl;

25 R⁸, R⁹, R¹⁶ and R¹⁷ independently represent hydrogen, -C₁₋₆alkyl, -C₃₋₆cycloalkyl, -C₁₋₃ alkylC₃₋₆ cycloalkyl, -C₂₋₆alkenyl or NR⁸R⁹ or NR¹⁶R¹⁷ together represents morpholinyl, pyrrolidinyl, piperidinyl, piperazinyl or piperazinyl N-substituted by -C₁₋₆ alkyl, -COphenyl or -SO₂methyl;

R¹⁰, R¹¹, R¹², R¹³, R¹⁵, R¹⁸, R²⁰ and R²¹ independently represent hydrogen or -C₁₋₆alkyl;

R^{14} , R^{19} and R^{22} independently represent hydrogen, $-C_{1-6}\text{alkyl}$, $-C_{3-6}\text{ cycloalkyl}$ or $-(CH_2)_x\text{Ar}^4$ or $NR^{14}R^{18}$ or $NR^{15}R^{22}$ together represents morpholinyl, pyrrolidinyl, piperidinyl, piperazinyl or $N-C_{1-6}\text{alkylpiperazinyl}$;

- 5 Ar^1 represents phenyl or a 5 or 6 membered heterocyclic aromatic ring containing 1 to 3 heteroatoms selected from O, N and S optionally substituted by one or more halogen, $C_{1-6}\text{alkyl}$, hydroxy, $-OC_{1-6}\text{alkyl}$, CF_3 , nitro, $-\text{Ar}^2$ or $-\text{OAr}^2$ groups;
- 10 Ar^2 represents phenyl optionally substituted by one or more halogen, $-C_{1-6}\text{alkyl}$, hydroxy, $-OC_{1-6}\text{alkyl}$, $-CF_3$ or nitro groups;
- 15 Ar^3 represents phenyl, a 5 or 6 membered heterocyclic aromatic ring containing 1 to 3 heteroatoms selected from O, N or S, or such a group fused to a benzene ring, optionally substituted by one or more $-\text{CO}(CH_2)_g\text{Ar}^4$, $-(CH_2)_y\text{Ar}^4$, $-(CH_2)_y\text{COAr}^4$, $-(CO)_aC_{1-6}$ alkyl, $-(CO)_aC_{2-6}$ alkenyl, $-(CO)_aC_{2-6}$ alkynyl, $-(CO)_aC_{3-8}\text{cycloalkyl}$, $-(CO)_aC_{1-6}\text{haloalkyl}$, halogen, $-\text{COCH}_2\text{CN}$, $-(CH_2)_b\text{NR}^{16}R^{17}$, $-(CH_2)_b\text{NHC(=NH)NH}_2$, $-\text{CYNR}^{16}(CO)_aR^{17}$, $-(CH_2)_b\text{NR}^{15}\text{COR}^{19}$, $-(CH_2)_b\text{CONR}^{15}R^{22}$, $-(CH_2)_b\text{NR}^{15}\text{CONR}^{15}R^{22}$, $-(CH_2)_b\text{CONR}^{15}(CH_2)_j\text{NR}^{15}R^{22}$, $-(CH_2)_b\text{SO}_2\text{NR}^{15}R^{22}$, $-(CH_2)_b\text{SO}_2\text{NR}^{15}\text{COAr}^2$, $-(CH_2)_b\text{NR}^{15}\text{SO}_2\text{R}^{19}$, $-\text{SO}_2\text{R}^{19}$, $-\text{SOR}^{19}$, $-(CH_2)_z\text{OH}$, $-\text{COOR}^{15}$, $-\text{CHO}$, $-OC_{1-10}\text{alkyl}$, $-\text{O}(CH_2)_j\text{NR}^{15}R^{22}$, $-\text{O}(CH_2)_j\text{NHC(=NH)NH}_2$, $-\text{O}(CH_2)_b\text{CONR}^{16}R^{17}$, $-\text{O}(CH_2)_k\text{COOR}^{15}$, $-\text{O}(CH_2)_j\text{OAr}^2$, $-\text{O}(CH_2)_b\text{Ar}^2$, 3-phenyl-2-pyrazolin-5-one or 4,5-dihydro-3(2H)-pyridazinone groups;
- 25 Ar^4 represents phenyl or a 5 or 6 membered heterocyclic aromatic ring containing 1 to 3 heteroatoms selected from O, N and S optionally substituted by one or more halogen, $-C_{1-6}\text{alkyl}$, hydroxy, $-OC_{1-6}\text{alkyl}$, $-CF_3$, nitro or $-\text{CONH}_2$ groups;

X and Y independently represent O or S;

a, f, k, s and n independently represent 0 or 1;

5 b, c, r, x, y and z independently represent an integer 0 to 2;

d, g and u independently represent 1 or 2;

e, h, q and w independently represent an integer 1 to 3;

j and p independently represent an integer 2 to 4;

m independently represents an integer 0 to 4;

10 t independently represents an integer 0 to 3;

and salts and solvates thereof.

Examples of 5 or 6 membered heterocyclic aromatic rings that Ar¹, Ar³ and Ar⁴ may represent include pyrimidine, pyridine, furan, imidazole, thiophene, pyrrole,

15 thiazole, oxazole, isoxazole, 1,3,4-thiadiazole, 1,2,3-thiadiazole, 1,2,4-thiadiazole, 1,2,4-oxadiazole and pyrazole.

Specific examples of 5 or 6 membered heterocyclic aromatic rings that Ar¹ may represent include pyrimidine, pyridine, furan, 1,2,4-thiadiazole and pyrrole.

20

Specific examples of 5 or 6 membered heterocyclic aromatic rings that Ar³ may represent include thiazole and pyridine. Phenyl fused to a benzene ring represents naphthyl. An example of a 5 or 6 membered heterocyclic aromatic ring fused to a benzene ring that Ar³ may represent includes benzofuran.

25

Specific examples of 5 or 6 membered heterocyclic aromatic rings that Ar⁴ may represent include 1,3,4-thiadiazole, 1,2,3-thiadiazole, 1,2,4-oxadiazole and pyrazole.

We prefer R¹ and R² to be defined such that NR¹R² together represent piperidinyl, piperazinyl, thiomorpholinyl, morpholinyl or 1,2,3,4-tetrahydroisoquinoline optionally substituted by a -(CO)_n(CH₂)_rAr¹, -(CO)_nC₁₋₆alkyl, -(CH₂)_tCONR⁸R⁹, -NR¹⁰(CO)_n(CH₂)_rAr¹, 5 -NR¹⁰(CO)_nC₁₋₃ alkylC₃₋₆ cycloalkyl, -NR¹⁰(CO)_nC₁₋₆ alkylidiC₃₋₆ cycloalkyl, -(CH₂)_rOC₁₋₆ alkyl, -(CH₂)_rO(CH₂)_pOH, piperidin-1-yl, -(CH₂)_rOH or -CONR¹⁰(CH₂)_rAr¹ group.

We particularly prefer R¹ and R² to be defined such that NR¹R² together 10 represents morpholinyl or piperazinyl optionally N-substituted by -(CO)_nC₁₋₆ alkyl (especially -COCH₃), piperazinyl N-substituted by -(CO)_n(CH₂)_rAr¹ (especially -COPhenyl and -(CO)2-furanyl), piperidinyl substituted by -NR¹⁰(CO)_n(CH₂)_rAr¹ (especially -NHCOCH₂phenyl) or piperidinyl substituted by -(CH₂)_tCONR⁸R⁹ (especially -CONH₂).

15 We prefer R³ to represent -(CH₂)_c-2,4-imidazolidinedione-3-yl, -(CH₂)_c-thioxanthen-9-one-3-yl, -(CH₂)_cAr³, -O(CH₂)_cAr³, -(CH₂)_dOAr³ or -(CH₂)_ddibenzofuran, particularly -OCH₂Ar³, -CH₂OAr³ or dibenzofuran, especially -CH₂OAr³ or dibenzofuran.
20 When R³ represents -(CH₂)_ddibenzofuran (particularly dibenzofuran), we prefer it to represent -(CH₂)_d-2-dibenzofuran (particularly 2-dibenzofuran).

When R³ represents -(CH₂)_c-2,4-imidazolidinedione, we prefer it to represent 25 -(CH₂)_c-(2,4-imidazolidinedione-3-yl) (particularly -CH₂-2,4-imidazolidinedione-3-yl).

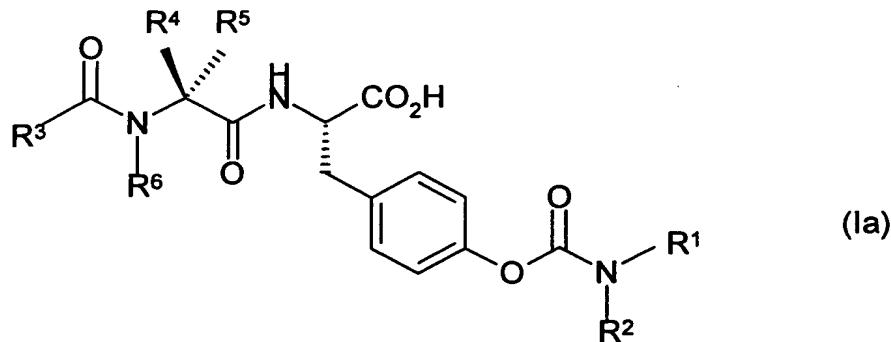
When R³ represents -(CH₂)_c-thioxanthen-9-one, we prefer it to represent -(CH₂)_c-(thioxanthen-9-one-3-yl) (particularly -CH₂-thioxanthen-9-one-3-yl).

We most especially prefer R³ to represent -CH₂OAr³.

We prefer R⁴ to represent -C₁₋₆ alkyl, R⁵ to represent hydrogen or for R⁴R⁵, together with the carbon to which they are attached, to form a cyclohexyl ring, and for R⁶ to represent hydrogen or methyl (particularly hydrogen).

5 We particularly prefer R⁴ to represent -C₁₋₆ alkyl, and for R⁵ and R⁶ to represent hydrogen. We especially prefer R⁴ to represent -CH₂CHMe₂ and for R⁵ and R⁶ to represent hydrogen.

10 We particularly prefer R⁴ and R⁵ to have the stereochemical orientation shown in formula (la):



15 We prefer R⁷ to represent -(CH₂)_vAr² or -(CH₂)_wNR¹²COC₁₋₆ alkyl.

We especially prefer R⁸ and R⁹ each to represent hydrogen or for NR⁸R⁹ together to represent piperidinyl or pyrrolidinyl, particularly piperidinyl.

We prefer R¹⁰ to represent hydrogen or methyl, particularly hydrogen.

20 We prefer R¹¹ to represent hydrogen or methyl, particularly hydrogen.

We prefer R¹² to represent hydrogen or methyl, particularly hydrogen.

We prefer R¹³ to represent hydrogen or methyl, particularly hydrogen.

We prefer R¹⁴ to represent hydrogen or methyl, particularly hydrogen.

We prefer R¹⁵ to represent hydrogen or -C₁₋₆ alkyl, particularly hydrogen.

We prefer R¹⁶ to represent hydrogen, -C₁₋₄ alkyl or -C₂₋₄ alkenyl, particularly hydrogen or propenyl.

We prefer R¹⁷ to represent hydrogen, -C₁₋₄ alkyl or -C₂₋₄ alkenyl, particularly hydrogen, methyl or propenyl.

5 We prefer R¹⁸ to represent hydrogen or methyl, particularly hydrogen.

We prefer R¹⁹ to represent hydrogen or -C₁₋₆ alkyl, particularly -C₁₋₆ alkyl, especially methyl.

We prefer R²⁰ to represent hydrogen or methyl, particularly hydrogen.

10 We prefer R²¹ to represent hydrogen or methyl, particularly hydrogen.

We prefer R²² to represent hydrogen, -C₁₋₄ alkyl or -(CH₂)_xAr⁴ or for NR¹⁵R²² together to represent piperidinyl, pyrrolidinyl or morpholinyl.

We especially prefer R¹⁵ and R²² to be defined such that NR¹⁵R²² together represents piperidinyl.

15

We prefer Ar¹ to represent furan, pyrimidine or phenyl optionally substituted by halogen (eg chlorine or fluorine) or -OC₁₋₆ alkyl.

We prefer Ar² to represent unsubstituted phenyl.

20

We prefer Ar³ to represent phenyl, naphthyl or benzofuran optionally substituted by one or more -(CH₂)_yCOAr⁴, -COOR¹⁵, -(CH₂)_bSO₂NR¹⁵R²², -(CH₂)_bNR¹⁵SO₂R¹⁹, -SO₂R¹⁹, (CO)_aC₂₋₆ alkenyl, -(CO)_aC₁₋₆ alkyl, -(CO)_aC₃₋₈cycloalkyl, halogen, -(CH₂)_bCONR¹⁵R²², 3-phenyl-2-pyrazolin-5-one-2-yl or 4,5-dihydro-3(2H)-pyridazinone-6-yl groups. We particularly prefer Ar³ to represent phenyl or naphthyl optionally substituted by -(CO)_aC₁₋₆ alkyl, -(CO)_aC₃₋₈ cycloalkyl, halogen, -(CH₂)_yCOAr⁴ or -(CH₂)_bCONR¹⁵R²².

We most particularly prefer Ar³ to represent phenyl substituted by n-propyl, tertiary butyl, cyclohexyl, iodine, -COphenyl or COPiperidin-1-yl or naphthyl substituted by COPiperidin-1-yl.

5 We prefer Ar⁴ to represent phenyl or furan optionally substituted by halogen, especially unsubstituted phenyl or furan.

We prefer e to represent 1 or 2.

We prefer n to represent 0 or 1.

10 We prefer r to represent 0 or 1, particularly 1.

We prefer p to represent 2.

We prefer t to represent 0, 1 or 3, particularly 0 or 1, especially 0.

We prefer h to represent 1 or 2, particularly 2.

We prefer d to represent 1.

15 We prefer m to represent 0 or 1, particularly 1.

We prefer c to represent 0 or 1, particularly 1.

We prefer f to represent 1.

We prefer q to represent 1 or 2, particularly 1.

We prefer u to represent 1.

20 We prefer w to represent 1 or 2, particularly 1.

We prefer x to represent 0 or 1, particularly 1.

We prefer a to represent 0.

We prefer y to represent 0 or 1, particularly 0.

We prefer b to represent 0 or 1, particularly 0.

25 We prefer j to represent 2 or 3, particularly 2.

We prefer z to represent 0 or 1, particularly 0.

We prefer k to represent 1.

We prefer s to represent 0.

We prefer g to represent 1.

We prefer X to represent oxygen.

We prefer Y to represent oxygen.

The most preferred compounds of formula (I) are:

5 (2S)-2-[(2S)-2-{[2-(2-Iodophenoxy)acetyl]amino}-4-methyl pentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid;

10 (2S)-2-{[(2S)-2-{[2-[2-(Tert-butyl)phenoxy]acetyl]amino}-4-methyl pentanoyl]amino}-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid;

15 (2S)-3-(4-{[(4-Acetyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-{[(2S)-2-{[2-[2-(tert-butyl)phenoxy]acetyl]amino}-4-methylpentanoyl]amino}propanoic acid;

20 (2S)-2-[(2S)-2-{[2-(2-Cyclohexylphenoxy)acetyl]amino}-4-methyl pentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid;

25 (2S)-2-[(2S)-4-Methyl-2-{[2-(2-methylphenoxy)acetyl]amino} pentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid;

 (2S)-2-{[(2S)-2-{[2-[2-(Tert-butyl)phenoxy]acetyl]amino}-4-methyl pentanoyl]amino}-3-{4-[(4-[(2-phenylacetyl)amino]-1-piperidinyl]carbonyl) oxy]phenyl}propanoic acid;

 (2S)-3-(4-{[(4-Acetyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-[(2S)-4-methyl-2-{[2-(2-methylphenoxy)acetyl]amino}pentanoyl]amino]propanoic acid;

 (2S)-3-(4-{[(4-Benzoyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-{[(2S)-2-{[2-[2-(tert-butyl)phenoxy]acetyl]amino}-4-methylpentanoyl]amino}propanoic acid;

 (2S)-3-(4-{[(4-Acetyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl]amino)propanoic acid;

 (2S)-2-{[(2S)-2-{[2-[2-(Tert-butyl)phenoxy]acetyl]amino}-4-methyl pentanoyl]amino}-3-[4-({[4-(2-furoyl)-1-piperazinyl]carbonyl)oxy}phenyl]propanoic acid;

(2S)-2-((2S)-2-[(Dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)-3-[4-({[4-(2-furoyl)-1-piperazinyl]carbonyl}oxy)phenyl]propanoic acid;

(2S)-3-(4-{[(4-Benzoyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-[(2S)-4-methyl-2-{[2-(2-methylphenoxy)acetyl]amino}pentanoyl)amino]propanoic acid;

(2S)-3-[4-({[4-(Aminocarbonyl)-1-piperidinyl]carbonyl}oxy)phenyl]-2-[(2S)-4-methyl-2-{[2-(2-methylphenoxy)acetyl]amino}pentanoyl)amino] propanoic acid;

(2S)-3-(4-{[(4-Benzoyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)propanoic acid;

10 and salts or solvates thereof.

(2S)-3-[4-({[4-(Aminocarbonyl)-1-piperidinyl]carbonyl}oxy)phenyl]-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino) propanoic acid;

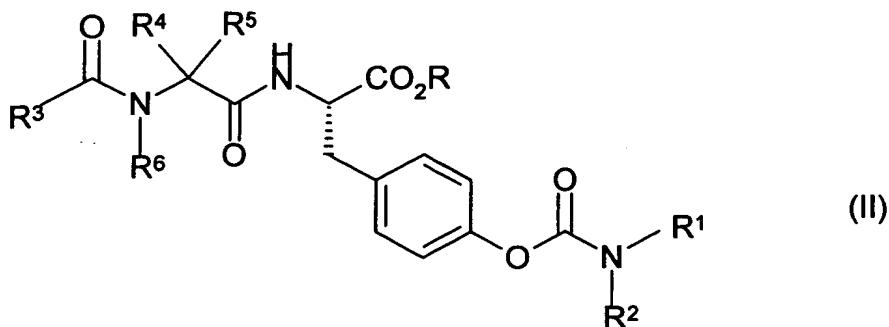
and salts or solvates thereof.

15

The compounds of formula (I) and salts and solvates thereof may be prepared by the methodology described hereinafter, constituting a further aspect of this invention.

20 A process according to the invention for preparing a compound of formula (I) comprises:

(a) hydrolysis of a carboxylic acid ester of formula (II)



wherein R^1 , R^2 , R^3 , R^4 , R^5 and R^6 are as defined above and R is a group capable of forming a carboxylic acid ester; or

5

(b) deprotecting a compound of formula (I) which is protected.

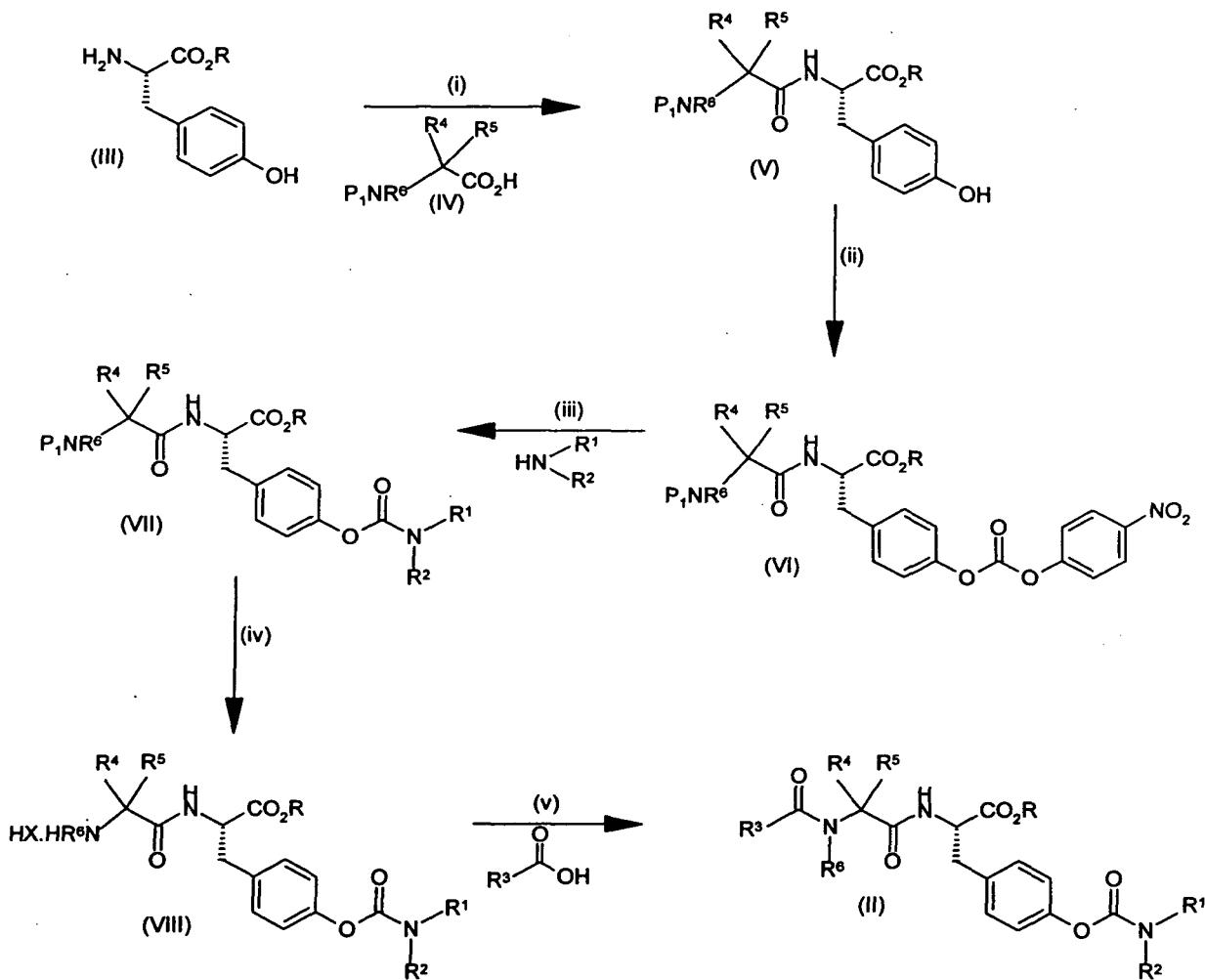
In process (a) an example of a suitable R group is a C_{1-6} alkyl group such as methyl or t-butyl. Hydrolysis may either occur via an acidic process e.g. involving trifluoroacetic acid and water or via an alkaline route e.g. utilising sodium hydroxide and methanol.

In an alternative solid phase reaction, R may represent a solid support functionalised with available hydroxy groups. Examples of solid supports include resins such as polystyrene resins wherein phenyl rings are provided with hydroxy groups via linkers. An example of a hydroxy functionalised linker is - $CH_2O(4\text{-hydroxymethyl-phenyl})$ (Wang Resin).

In process (b) examples of protecting groups and the means for their removal can be found in T. W. Greene 'Protective Groups in Organic Synthesis' (J. Wiley and Sons, 1991). Suitable amine protecting groups include sulphonyl (e.g. tosyl), acyl (e.g. benzyloxycarbonyl or t-butoxycarbonyl) and arylalkyl (e.g. benzyl), which may be removed by hydrolysis or hydrogenolysis as appropriate.

Compounds of formula (II) may be prepared following Scheme 1;

Scheme 1



Step (i)

In this Scheme we prefer R to represent methyl.

Compounds of formula (III) and (IV) may be reacted under conventional conditions for preparation of an amide. Desirably a coupling agent eg WSCDI with or without HOBT in an inert solvent such as MeCN or DMF is used. P₁ is an amine protecting group such as one described previously under process (b). In
5 this Scheme we prefer P₁ to represent Boc.

Step (ii)

The conversion of formula (V) to (VI) is suitably carried out with p-nitrophenylchloroformate under conventional conditions eg. in the presence of
10 an organic base, eg. pyridine and an inert organic solvent such as DCM.

Step (iii)

15 This reaction may be performed by combination of the reagents in a suitable solvent, such as DCM in the presence of an organic base such as DIPEA.

Step (iv)

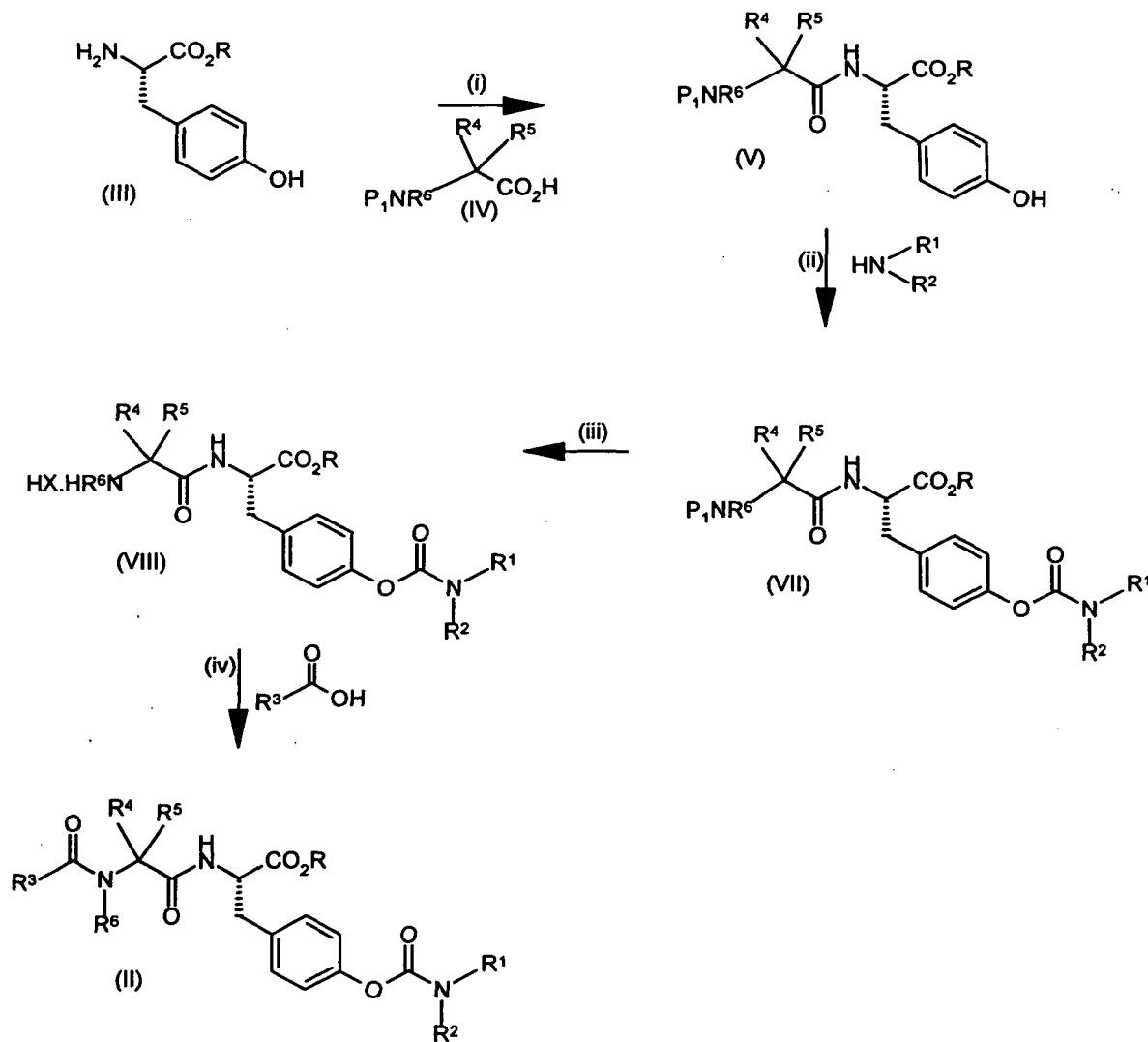
20 This deprotection step may be performed under conventional conditions. When P₁ represents Boc, it may be removed by treatment with acid e.g. a hydrohalic acid (HX) such as HCl.

Step (v)

25 A condensation reaction of formula (VIII) with the compound of formula R³CO₂H may be performed under conditions similar to those described above for step (i).

An alternative process for preparation of compounds of formula (II) is given in Scheme 2 below:

Scheme 2



Step (i)

In this Scheme we prefer R to represent t-Bu. The reaction conditions for this step are analogous to those for Scheme 1 step (i).

5 In compounds of formula (IV) in this Scheme we prefer P¹ to represent Cbz.

Step (ii)

This process comprises a two stage reaction, consisting of (a) treatment with a carboxyl donor such as (Cl₃CO)₂CO typically in the presence of an organic base such as DIPEA and a suitable solvent, such as THF or DCM followed by (b) conversion to the carbamate by treatment with R¹R²NH in a process analogous to that described previously in Scheme 1 step (iii).

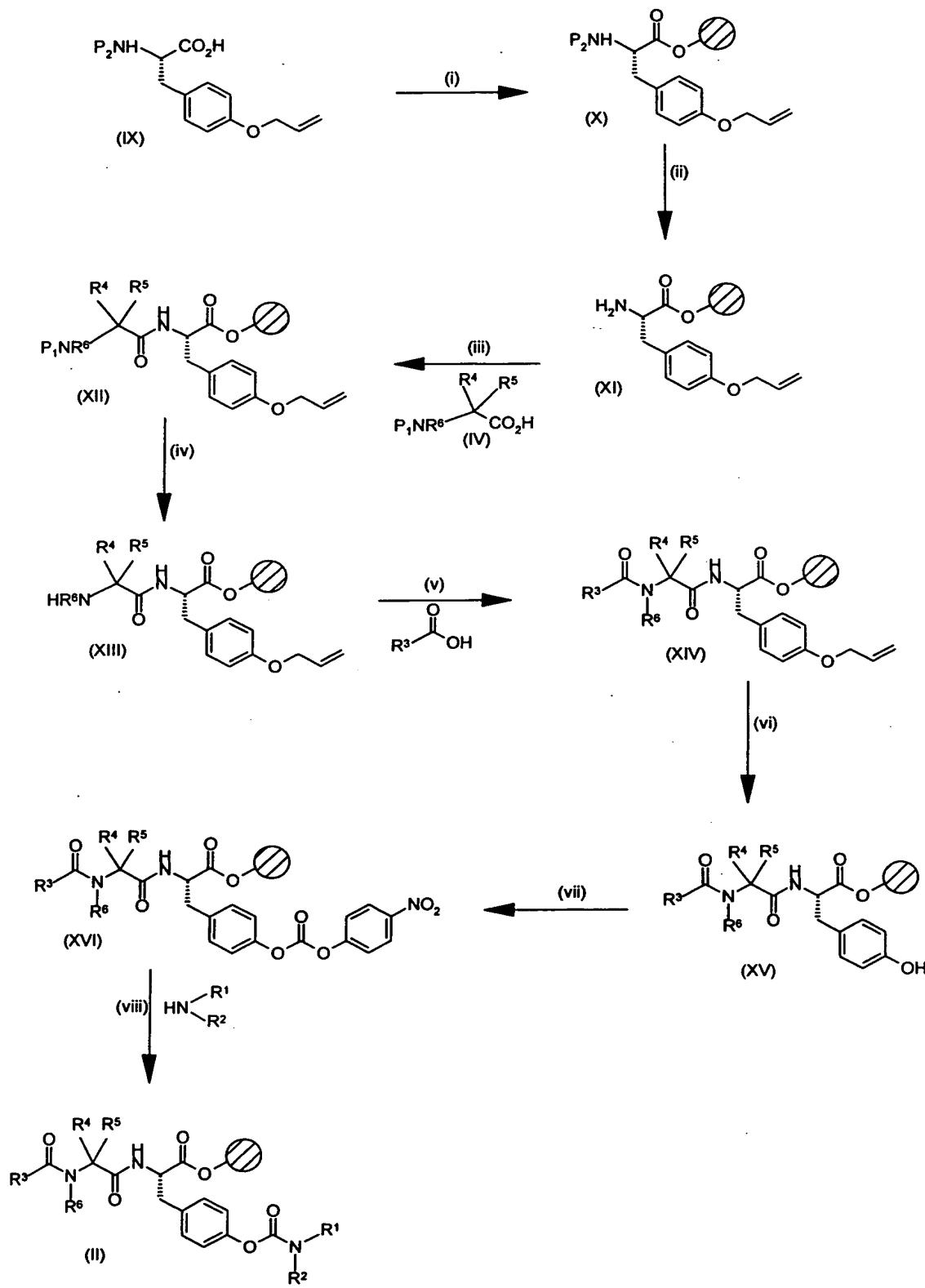
15 Step (iii)

This deprotection reaction can be performed under conventional conditions. When P₁ represents Cbz, deprotection may be achieved by hydrogenolysis e.g. by treatment with ammonium formate in the presence of Pd/C in a solvent such as ethanol. The reaction may be worked up with acid, such as a hydrohalic acid to give the product as a hydrohalic acid salt (e.g. the HCl salt).

Step (iv)

25 This process is analogous to Scheme 1, step (v).

An alternative process for preparation of compounds of formula (II) is given in Scheme 3 below:

Scheme 3

Step (i)

P₂ is an amine protecting group such as one described previously and in this Scheme we prefer P₂ to represent Fmoc.

5

A compound of formula (IX) may be reacted onto a suitable solid phase, such as a hydroxy functionalised polystyrene resin (e.g. Wang resin) in the presence of 2,6-dichlorobenzoyl chloride, pyridine and a suitable solvent, such as DMF.

10

Step (ii)

Removal of N-protecting group P₂ may be achieved under conventional conditions; e.g. when P₂ represents Fmoc, by treatment with an organic base such as piperidine in a suitable solvent, such as DMF

15

Step (iii)

20 In this Scheme, P₁ may suitably represent Fmoc. Reaction of a compound of formula (XI) with the compound of formula (IV) to produce an amide, may be performed in the presence of a coupling agent, such as PyBop, an organic base, such as DIPEA and a suitable solvent, such as DMF.

Step (iv)

25 This de-protection reaction may be performed under conventional conditions e.g. when P₁ represents Fmoc, under conditions analogous to those described above for step (ii).

Step (v)

5 A condensation reaction of formula (XIII) with the compound of formula R^3CO_2H may be performed in the presence of a suitable coupling agent, such as PyBop, an organic base, such as DIPEA and a suitable solvent, such as DMF.

Step (vi)

10 This step comprises an alkenyl chain cleavage reaction on the compound of formula (XIV) to produce a compound of formula (XV), eg. by the treatment with $Pd(PH_3)_4$ and $PhSiH_3$ in the presence of a suitable solvent, such as DCM.

Step (vii)

15

The conversion of a compound of formula (XV) to a compound of formula (XVI) is suitably performed by treatment with p-nitrophenyl chloroformate, under conventional conditions, in the presence of an organic base, such as DIPEA and an inert organic solvent, such as THF and/or DCM.

20

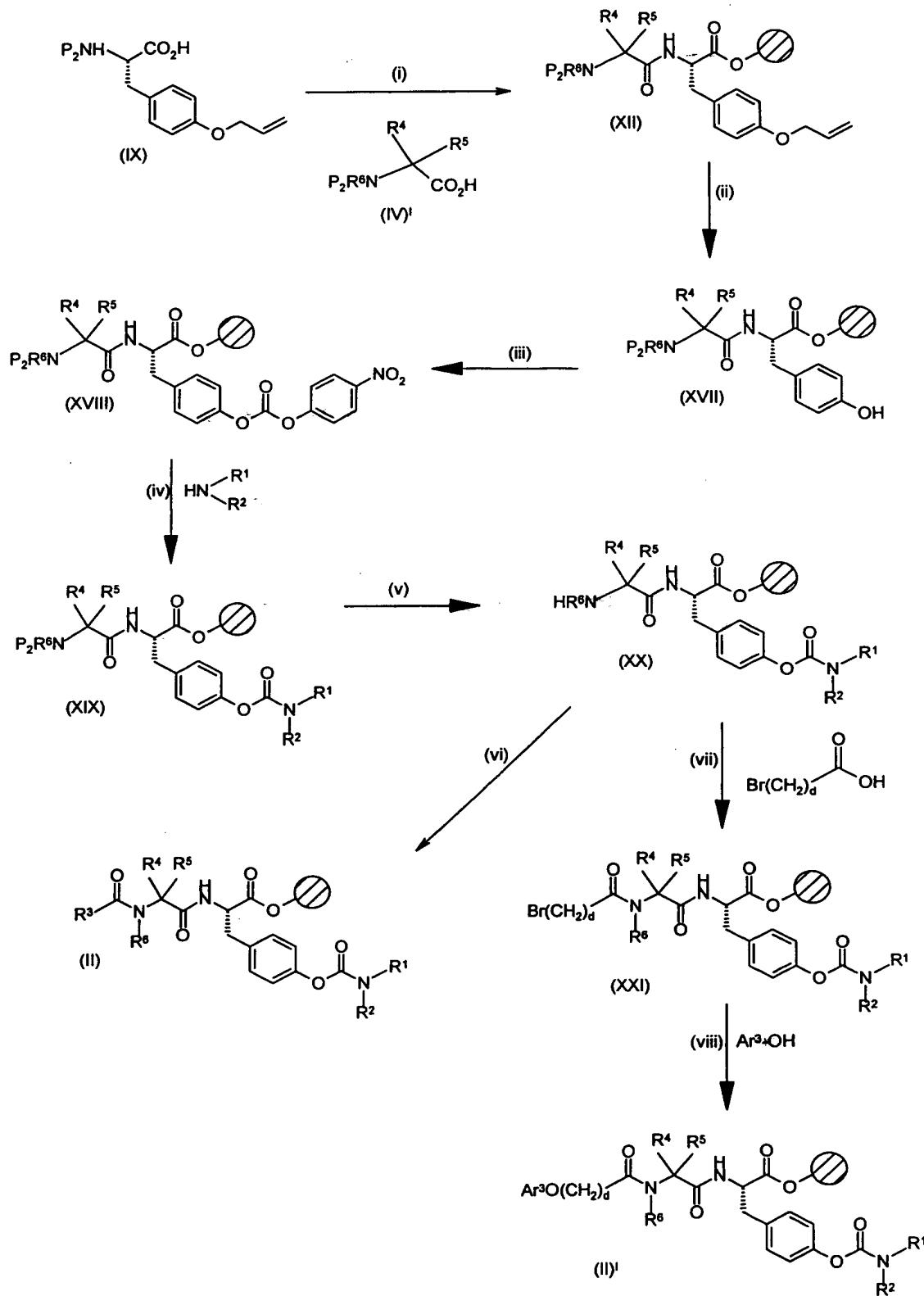
Step (viii)

25

This reaction may be performed by combination of the reagents in the presence of an organic base, such as DIPEA and suitable solvents, such as DCM and/or THF.

An alternative process for preparation of compounds of formula (II) is given in Scheme 4 below:

Scheme 4



Step (i)

In this Scheme we prefer P₂ to represent Fmoc.

5 This conversion may be achieved following processes analogous to those of Scheme 3 steps (i) to (iii).

Step (ii)

10 An alkenyl chain cleavage reaction may be performed by a process analogous to Scheme 3 step (vi).

Step (iii)

15 A p-nitrophenyl carbonate formation reaction, may be performed with reaction conditions analogous to Scheme 3 step (vii).

Step (iv)

20 The conversion of formula (XVIII) to (XIX) can be performed by a reaction analogous to Scheme 3 step (viii).

Step (v)

25 This de-protection reaction may be performed using an analogous process to Scheme 3 step (ii).

Step (vi)

5 The conversion of formula (XX) to (II) can be performed by a condensation reaction in the presence of a suitable acid, employing a suitable coupling agent, such as PyBop, an organic base, such as DIPEA and a solvent, such as DMF.

Compounds of formula (II) in which R³ represents -(CH₂)_dOAr³ may alternatively be prepared from compounds of formula (XX) following steps (vii) and (viii):

10 Step (vii)

The conversion of formula (XX) to (XXI) can be performed by a condensation reaction in the presence of a haloalkanoic acid (such as the bromo derivative), employing a suitable coupling agent, such as DIC and a solvent, such as DMF.

15

Step (viii)

20 In this step, the reaction of a compound of formula (XXI) with a compound of formula Ar³-OH group may be undertaken in the presence of potassium carbonate, sodium iodide and a suitable solvent, such as DMF.

Compounds of formula III, IV, IV¹, HNR¹R², R³COOH, IX, Br(CH₂)_eCOOH and Ar³-OH are either known or may be prepared by known methods.

25 Compounds of the invention may be tested for *in vitro* and *in vivo* biological activity in accordance with the following assays.

(1) Jurkat J6/VCAM-1 Adhesion Assay

This assay was used to investigate the interaction of the integrin VLA-4, expressed on the Jurkat J6 (human lymphoblast cell line) cell membrane with VCAM-1. Polystyrene 96-well microtitre plates were coated with human immunoglobulin G (IgG; Sigma Chemicals, UK, Product No. I4506) at a concentration of 0.05mg ml^{-1} in bicarbonate buffer (36mM NaHCO_3 and 22mM Na_2CO_3 , prepared in Dulbecco's phosphate buffered saline at pH 9.8 (PBS); Sigma Chemicals, UK, Product No. 14190-094) for 2 hours at 37°C . This solution was then aspirated and the plates were washed twice with PBS.

10

VCAM-1 was prepared by cloning its constituent seven domains into a Drosophila expression system with a zz (Protein A) tag. This zzVCAM-1 was then expressed from *Drosophila melanogaster* S2 cell culture, induced with copper. Protease inhibitors were added and the culture supernatant was clarified either by filtration through a $0.2\mu\text{m}$ filter or by centrifugation. The zzVCAM-1 was then purified from this clarified medium using an IgG agarose column, equilibrated with either 20mM sodium phosphate pH 7.2 alone or in the presence of 0.5M sodium chloride. Elution of zzVCAM-1 from the column was mediated using 3M ammonium thiocyanate, which was subsequently removed using a G25 desalting column, equilibrated with 20mM sodium phosphate, pH 7.2. The purified zzVCAM-1 was then concentrated to a small volume (Amicon stirred cell concentrators) until a concentration of 62.5ng ml^{-1} was obtained, calculated using the extinction coefficient value.

25

This solution of zzVCAM-1 was then incubated overnight at 4°C in the IgG coated microtitre plates with 3% bovine serum albumin (BSA) in PBS, followed by aspiration and two further washes with PBS. A concentration of the Jurkat J6 cells (6×10^6 cells ml^{-1}), grown in cell media RPMI 1640 (HyClone Ltd, Product No. B-9106-L) supplemented with 10% heat inactivated foetal calf serum (FCS;

Gibco BRL, Product No. 10099-075) and 2mM L-glutamine, were labelled with 10 μ M of the fluorescent dye, 2', 7'-bis(2-carboxyethyl)-5-(e6)-carboxyfluorescein acetoxyethyl ester (BCECF-AM; Molecular Probes Inc, Product No. B-1150) at 37°C for 10 minutes. The excess dye was then removed by centrifugation at 5 500xg for 5 minutes and the cells were resuspended at a concentration of 1.2 x 10⁷ cells ml⁻¹ in Hank's balanced salt solution (HBSS; Gibco BRL, Product No. 14190-094).

Equal volumes of compounds (dissolved in an appropriate solvent and diluted in 10 HBSS containing 1mM MnCl₂) and the labelled Jurkat J6 cells, were added to the VCAM-1 coated plates and adhesion was allowed to proceed for 30 minutes at 37°C. Non, or loosely adhered cells, were removed by inversion of the plate and blotted with tissue paper. Two washes with PBS and further blotting were then performed, before the addition of 2% detergent (Triton-X®; Sigma 15 Chemicals UK, Product No. X100). Counting was undertaken in a Wallac Viktor™ Fluorimeter, where low fluorescence values were indicative of compounds which had inhibited adhesion. All samples were assayed in singlicate and the following four parameter curve fit, shown by Equation (I) was applied:

Equation (I)

$$y = \frac{a - d}{1 + (\frac{x}{c})^b} + d$$

20 Where a is the minimum, b is the Hill slope, c is the IC₅₀ and d is the maximum. (Maximum and minimum values are those compared to adhesion in the absence of compound and in the presence of the dipotassium salt of 2mM EDTA; Sigma Chemicals, UK, Product No. ED2P). Data is presented as the mean pIC₅₀ with 25 the standard error of the mean of n experiments.

(2) CD4/VCAM-1 Co-stimulation of T-Lymphocyte Proliferation

CD4+ T-cells were purified from peripheral blood mononuclear cells by negative selection with anti-CD14, CD19, CD16 and HLA.DR antibodies and Dynal beads. Flat bottomed 96-well tissue culture plates were coated with 1 μ g ml $^{-1}$

5 anti-CD3 antibody (OKT3), washed and incubated with human IgG and

zzVCAM-1 fusion proteins. The CD4+ T cells (prepared in RPMI-1640 medium supplemented with 10% FCS, penicillin or streptomycin and L-glutamine) were added to the coated plates (1×10^5 cells well $^{-1}$) and incubated in the presence or absence of various doses of compound or blocking antibodies for 4 days.

10 Radiolabelled thymidine [3 H] was added for the final 6 hours of incubation and the cells were then harvested using a Skatron plate harvester. Incorporation of the [3 H] label was measured as an indicator of T cell proliferation using a β plate counter. Compounds were assayed in triplicate and data was collected in an analogous procedure to that described for Assay (1).

15

(3) Inhibition of Eosinophil Infiltration and Hyper-Reactivity in the Guinea Pig

In a method based on that described by Danahay *et al.*, 1997, ovalbumin sensitised guinea pigs were dosed with mepyramine (30mg kg $^{-1}$ ip) to protect against anaphylactic bronchospasm. Test compounds, dissolved in 0.9% saline,

20 were given by the inhaled route (30 minutes breathing of an aerosol of the compound) or the intra-tracheal route, 30 minutes before and 6 hours after ovalbumin challenge (10 minutes breathing of an aerosol generated from a 0.5% solution of ovalbumin). Hyper-reactivity of the airways to the thromboxane mimetic U46619, was measured 24 hours after ovalbumin challenge in unrestrained animals using a whole body plethysmograph (Buxco Ltd., USA). The

25 guinea pigs were then sacrificed and the lungs lavaged. Total and differential leukocyte counts were then obtained for the bronchoalveolar lavage fluid and the percentage reduction in eosinophil accumulation determined (Sanjar *et al.*, 1992). Dexamethasone (1mg i.t) was used as a positive control. Data was

presented as the minimum dose that is maximally effective in attenuating eosinophil accumulation or airway hyperreactivity, together with the maximum inhibitory effect expressed as a percentage of the vehicle control response.

5 Examples of disease states in which the compounds of the invention have potentially beneficial anti-inflammatory effects include diseases of the respiratory tract such as bronchitis (including chronic bronchitis), asthma (including allergen-induced asthmatic reactions), chronic obstructive pulmonary disease (COPD) and rhinitis. Other relevant disease states include diseases of the 10 gastrointestinal tract such as intestinal inflammatory diseases including inflammatory bowel disease (e.g. Crohn's disease or ulcerative colitis) and intestinal inflammatory diseases secondary to radiation exposure or allergen exposure. Furthermore, compounds of the invention may be used to treat nephritis, skin diseases such as psoriasis, allergic dermatitis and 15 hypersensitivity reactions and diseases of the central nervous system which have an inflammatory component eg Alzheimer's disease, meningitis, multiple sclerosis and AIDS dementia.

Further examples of disease states in which compounds of the invention have 20 potentially beneficial effects include cardiovascular conditions such as atherosclerosis, peripheral vascular disease and idiopathic hypereosinophilic syndrome.

Compounds of the invention may be useful as immunosuppressive agents and 25 so have use in the treatment of auto-immune diseases such as allograft tissue rejection after transplantation, rheumatoid arthritis and diabetes.

Compounds of the invention may also be useful in inhibiting metastasis.

Diseases of principal interest include asthma, COPD and inflammatory diseases of the upper respiratory tract involving seasonal and perennial rhinitis.

It will be appreciated by those skilled in the art that reference herein to treatment extends to prophylaxis as well as the treatment of established conditions.

As mentioned above, compounds of formula (I) are useful as pharmaceuticals, in particular as anti-inflammatory agents.

10 There is thus provided as a further aspect of the invention a compound of formula (I) or a physiologically acceptable salt or solvate thereof for use as pharmaceuticals, particularly in the treatment of patients with inflammatory conditions.

15 According to another aspect of the invention, there is provided the use of a compound of formula (I) or a physiologically acceptable salt or solvate thereof for the manufacture of a medicament for the treatment of patients with inflammatory conditions.

20 In a further or alternative aspect there is provided a method for the treatment of a human or animal subject with an inflammatory condition, which method comprises administering to said human or animal subject an effective amount of a compound of formula (I) or a physiologically acceptable salt or solvate thereof.

25 The compounds according to the invention may be formulated for administration in any convenient way, and the invention therefore also includes within its scope pharmaceutical compositions for use in anti-inflammatory therapy, comprising a compound of formula (I) or a physiologically acceptable salt or solvate thereof

together, if desirable, with one or more physiologically acceptable diluents or carriers.

There is also provided a process for preparing such a pharmaceutical
5 formulation which comprises mixing the ingredients.

The compounds according to the invention may, for example, be formulated for oral, buccal, parenteral, topical or rectal administration, preferably for topical administration to the lung, eg. by aerosol or as a dry powder composition.

10 Tablets and capsules for oral administration may contain conventional excipients such as binding agents, for example syrup, acacia, gelatin, sorbitol, tragacanth, mucilage of starch, cellulose or polyvinyl pyrrolidone; fillers, for example, lactose, microcrystalline cellulose, sugar, maize- starch, calcium phosphate or sorbitol; lubricants, for example, magnesium stearate, stearic acid, talc, polyethylene glycol or silica; disintegrants, for example, potato starch, croscarmellose sodium or sodium starch glycollate; or wetting agents such as sodium lauryl sulphate. The tablets may be coated according to methods well known in the art. Oral liquid preparations may be in the form of, for example,
15 aqueous or oily suspensions, solutions, emulsions, syrups or elixirs, or may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations may contain conventional additives such as suspending agents, for example, sorbitol syrup, methyl cellulose, glucose/sugar syrup, gelatin, hydroxymethyl cellulose, carboxymethyl cellulose, aluminium
20 stearate gel or hydrogenated edible fats; emulsifying agents, for example, lecithin, sorbitan mono-oleate or acacia; non-aqueous vehicles (which may include edible oils), for example almond oil, fractionated coconut oil, oily esters, propylene glycol or ethyl alcohol; or preservatives, for example, methyl or propyl
25 p- hydroxybenzoates or sorbic acid. The preparations may also contain buffer

salts, flavouring, colouring and/or sweetening agents (e.g. mannitol) as appropriate.

For buccal administration the compositions may take the form of tablets or

5 lozenges formulated in conventional manner.

The compounds may also be formulated as suppositories, e.g. containing conventional suppository bases such as cocoa butter or other glycerides.

10 The compounds according to the invention may also be formulated for parenteral administration by bolus injection or continuous infusion and may be presented in unit dose form, for instance as ampoules, vials, small volume infusions or pre-filled syringes, or in multi-dose containers with an added preservative. The compositions may take such forms as solutions, suspensions,

15 or emulsions in aqueous or non-aqueous vehicles, and may contain formulatory agents such as anti-oxidants, buffers, antimicrobial agents and/or tonicity adjusting agents. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g. sterile, pyrogen-free water, before use.

20 The dry solid presentation may be prepared by filling a sterile powder aseptically into individual sterile containers or by filling a sterile solution aseptically into each container and freeze-drying.

By topical administration as used herein, we include administration by insufflation and inhalation. Examples of various types of preparation for topical 25 administration include ointments, creams, lotions, powders, pessaries, sprays, aerosols, capsules or cartridges for use in an inhaler or insufflator, solutions for nebulisation or drops (e.g. eye or nose drops).

Ointments and creams may, for example, be formulated with an aqueous or oily base with the addition of suitable thickening and/or gelling agents and/or solvents. Such bases may thus, for example, include water and/or an oil such as liquid paraffin or a vegetable oil such as arachis oil or castor oil or a solvent such as a polyethylene glycol. Thickening agents which may be used include soft paraffin, aluminium stearate, cetostearyl alcohol, polyethylene glycols, microcrystalline wax and beeswax.

5 Lotions may be formulated with an aqueous or oily base and will in general also contain one or more emulsifying agents, stabilising agents, dispersing agents, suspending agents or thickening agents.

10 Powders for external application may be formed with the aid of any suitable powder base, for example, talc, lactose or starch. Drops may be formulated with an aqueous or non-aqueous base also comprising one or more dispersing agents, solubilising agents or suspending agents.

15 Powder compositions for inhalation will preferably contain lactose. Spray compositions may be formulated, for example, as aqueous solutions or suspensions or as aerosols delivered from pressurised packs, with the use of a suitable propellant, e.g. dichlorodifluoromethane, trichlorofluoromethane, dichlorotetra-fluoroethane, 1,1,1,2,3,3-heptafluoropropane, 1,1,1,2-tetrafluoroethane, carbon dioxide or other suitable gas.

20 Intranasal sprays may be formulated with aqueous or non-aqueous vehicles with the addition of agents such as thickening agents, buffer salts or acid or alkali to adjust the pH, isotonicity adjusting agents or anti-oxidants.

Capsules and cartridges of for example gelatin, or blisters of for example laminated aluminium foil, for use in an inhaler or insufflator may be formulated containing a powder mix of a compound of the invention and a suitable powder base such as lactose or starch.

5

Solutions for inhalation by nebulisation may be formulated with an aqueous vehicle with the addition of agents such as acid or alkali, buffer salts, isotonicity adjusting agents or antimicrobials. They may be sterilised by filtration or heating in an autoclave, or presented as a non-sterile product.

10

The pharmaceutical compositions according to the invention may also be used in combination with other therapeutic agents, for example anti-inflammatory agents (such as corticosteroids (eg fluticasone propionate, beclomethasone dipropionate, mometasone furoate, triamcinolone acetonide or budesonide) or 15 NSAIDs (eg sodium cromoglycate, nedocromil sodium, PDE-4 inhibitors, leukotriene antagonists, iNOS inhibitors, tryptase and elastase inhibitors, beta-2 integrin antagonists and adenosine 2a agonists) or beta adrenergic agents (such as salmeterol, salbutamol, formoterol, fenoterol or terbutaline and salts thereof) or antiinfective agents (eg antibiotics, antivirals).

20

The invention thus provides, in a further aspect, a combination comprising a compound of formula (I) or a physiologically acceptable salt or solvate thereof together with another therapeutically active agent, for example an anti-inflammatory agent such as a corticosteroid, NSAID, beta adrenergic agent or 25 an anti-infective agent.

The combination referred to above may conveniently be presented for use in the form of a pharmaceutical formulation and thus pharmaceutical formulations

comprising a combination as defined above together with a physiologically acceptable diluent or carrier thereof represent a further aspect of the invention.

5 The individual components of such combinations may be administered either sequentially or simultaneously in separate or combined pharmaceutical formulations. Appropriate doses of known therapeutic agents will be readily appreciated by those skilled in the art.

10 Compounds of the invention may conveniently be administered in amounts of, for example, 0.01 to 500mg/kg body weight, preferably 0.01 to 100mg/kg body weight, 1 to 4 times daily. The precise dose will of course depend on the age and condition of the patient and the particular route of administration chosen.

15 The compounds of the invention have the advantage that they may be more efficacious, show greater selectivity, have fewer side effects, have a longer duration of action, be more bioavailable by the preferred route, show less systemic activity when administered by inhalation or have other more desirable properties than similar known compounds.

20 Certain intermediates are new and provide a further aspect of the invention.

The invention may be illustrated by reference to the following examples:

Examples

25 General Experimental Details

Where compounds were purified by "flash column chromatography on silica gel" this refers to the use of silica gel, 0.040 to 0.063mm mesh (e.g. Merck Art 9385), where column elution is accelerated by an applied pressure of nitrogen at up to

5 p.s.i. Where thin layer chromatography (TLC) has been used this refers to silica gel TLC using 5 x 10 cm silica gel plates (e.g. Polygram SIL G/UV₂₅₄).

Mass Spectroscopy

5 Mass Spectrometry (MS) was carried out using an HP5989A Engine Mass Spectrometer connected to a flow inject system (0.05M aqueous ammonium acetate/methanol (35:65) at a flow rate of 0.7 mL/min) with positive thermospray ionisation .

10

LC/MS System

The Liquid Chromatography Mass Spectrometry (LCMS) system used was as follows: -

A 3μm ABZ+PLUS, 3.3cm x 4.6mm internal diameter column eluting with solvents: A -0.01M Aqueous ammonium acetate + 0.1%v/v formic acid, and B -

15 95:5 acetonitrile/water + 0.05%v/v formic acid with a flow rate of 3mL/min. The following gradient protocol was used: 100% A for 0.7 mins; A+B mixtures, gradient profile 0 - 100% B over 3.7 mins; hold at 100% B for 0.9 mins; return to 0% B over 0.2 mins.

Positive and negative electrospray ionisation was employed.

20

Protection Measurement

The method for measuring the substitution of Fmoc-amino acid resins was as follows:-

To 10mg of resin was added 20% piperidine in DMF (1mL). After shaking for 30
25 mins at 20°C the resin was filtered. To 50μL of the filtrate was added 20% piperidine in DMF (0.95mL) and the absorbance of the solution was measured at 302nm using a UV spectrophotometer. Substitution was calculated using the following equation:-

Substitution (mmol/g) = (Absorbance x 2 x 10⁴) / (Extinction coefficient x weight in mg)

Intermediates

5 Intermediate 1: Methyl (2S)-2-{[(2S)-2-[(tert-butoxycarbonyl)amino]-4-methylpentanoyl]amino}-3-(4-hydroxyphenyl)propanoate

To a solution of N-(tert-butoxycarbonyl)-L-leucine (7g) in acetonitrile (100mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide-hydrochloride (5.9g) and 1-hydroxybenzotriazole (4.2g). After stirring for 30 mins at 20°C L-tyrosine methyl ester (5.5g) was added and stirring was continued for 18h. The mixture was concentrated *in vacuo* to ca. 10mL and the residue was partitioned between 1M hydrochloric acid (200mL) and ethyl acetate (100mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (100mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (100mL), water (2 x 100mL) and brine (50mL), dried over sodium sulphate and evaporated *in vacuo*. The residue was co-evaporated with chloroform to give the title compound as a white foam (11.3g, 98%). LCMS: R_t 3.11 min; *m/z* 409 (MH⁺).

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Intermediate 2: Methyl (2S)-2-{[(2S)-2-amino-4-methylpentanoyl]amino}-3-(4-hydroxyphenyl)propanoate hydrochloride

To a solution of Intermediate 1 (3.1g) in 1,4-dioxane (10mL) was added 4M hydrogen chloride in 1,4-dioxane (20mL). The solution was stirred for 2h at 20°C then evaporated *in vacuo*. The residue was co-evaporated with toluene (2 x 20mL) and ether (2 x 20mL) to give the title compound as a white solid (2.6g, 98%). LCMS: R_t 1.98 min; *m/z* 309 (MH⁺).

Intermediate 3: Methyl (2S)-3-(4-hydroxyphenyl)-2-((2S)-4-methyl-2-[(2-{[3-(1-piperidinylcarbonyl)-2-naphthyl]oxy}acetyl)amino]pentanoyl)amino)propanoate

To a suspension of Intermediate 44 (0.45g) in acetonitrile (20mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.31g) and 1-hydroxybenzotriazole (0.22g). After stirring for 30 mins at 20°C Intermediate 2 (0.5g) was added followed by diisopropylethylamine (0.28mL) and stirring was continued for 18h. The mixture was concentrated *in vacuo* and the residue was partitioned between 2M hydrochloric acid (50mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (30mL), water (2 x 30mL) and brine (20mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate/petroleum ether (2:1) to give the title compound as a white foam (0.6g, 69%). LCMS: R_t 3.42 min; *m/z* 604 (MH⁺).

Intermediate 4: Methyl (2S)-3-(4-hydroxyphenyl)-2-[(2S)-2-[(2-(2-iodophenoxy)acetyl)amino]-4-methylpentanoyl)amino]propanoate

This was similarly prepared from Intermediate 43 (0.81g) and Intermediate 2 (1.02g). The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) to give the title compound as a white foam (1.2g, 74%). LCMS: R_t 3.40 min; *m/z* 569 (MH⁺).

Intermediate 5: Methyl (2S)-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)-3-(4-hydroxyphenyl)propanoate

This was similarly prepared from Intermediate 45 (0.29g) and Intermediate 2 (0.5g). The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) to give the title compound as a white foam (0.66g, 97%). LCMS: R_t 3.55 min; *m/z* 503 (MH⁺).

Intermediate 6: Methyl (2S)-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)-3-(4-[(4-nitrophenoxy)carbonyloxy]phenyl)propanoate

To a solution of Intermediate 5 (0.59g) in dichloromethane (5mL), under a nitrogen atmosphere, was added 4-dimethylaminopyridine (0.18g). The mixture was cooled to 0-5°C and 4-nitrophenyl chloroformate (0.3g) was added. Stirring was continued for 18h allowing the reaction to warm to 20°C. The solution was diluted with chloroform (60mL) and washed with 1M hydrochloric acid (2 x 40mL) and water (40mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with cyclohexane/ethyl acetate (3:2) to give the title compound as a white foam (0.36g, 46%). LCMS: R_t 3.98 min; *m/z* 668 (MH⁺).

Intermediate 7: 4-[(2S)-2-((2S)-2-[(Tert-butoxycarbonyl)amino]-4-methylpentanoyl)amino]-3-methoxy-3-oxopropyl]phenyl 4-[(2-phenylacetyl)amino]-1-piperidinecarboxylate

To a solution of triphosgene (0.59g) in anhydrous dichloromethane (40mL), under a nitrogen atmosphere, was added a solution of Intermediate 1 (1.87g) in anhydrous dichloromethane (10mL) followed by diisopropylethylamine (1.2mL). After stirring for 3h at 20°C Intermediate 59 (1g) was added followed by diisopropylethylamine (0.8mL). Stirring was continued for 18h then the mixture was evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1 switching to 5:1) to give the title compound as a white solid (1.76g, 59%).

LCMS: R_t 3.42 min; *m/z* 651 [M-H]⁻.

Intermediate 8: 4-((2S)-2-[(2S)-2-Amino-4-methylpentanoyl]amino)-3-methoxy-3-oxopropyl]phenyl 4-[(2-phenylacetyl)amino]-1-piperidinecarboxylate hydrochloride

To a solution of Intermediate 7 (1.76g) in 1,4-dioxane (10mL) was added 4M hydrogen chloride in 1,4-dioxane (8mL). After stirring for 3h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a cream solid (1.59g, 100%). LCMS: R_t 2.50 min; m/z 553 (MH⁺).
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Intermediate 9: Methyl (2S)-2-{[(2S)-2-[(tert-butoxycarbonyl)amino]-4-methyl pentanoyl}amino)-3-(4-[(4-nitrophenoxy)carbonyloxy]phenyl)propanoate

To a solution of Intermediate 1 (0.41g) in dichloromethane (3mL), under a nitrogen atmosphere, was added pyridine (1mL). The mixture was cooled to 0-5°C and 4-nitrophenyl chloroformate (0.22g) was added. Stirring was continued for 18h allowing the reaction to warm to 20°C. The solution was diluted with dichloromethane (40mL) and washed with 1M hydrochloric acid (50mL). The aqueous phase was further extracted with dichloromethane (40mL) and the combined organic extracts were dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with petroleum ether/ethyl acetate (3:1 switching to 3:2) to give the title compound as a white solid (0.29g, 50%). LCMS: R_t 3.39 min; m/z 574 (MH⁺).
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Intermediate 10: 4-((2S)-2-{[(2S)-2-Amino-4-methylpentanoyl]amino}-3-methoxy-3-oxopropyl)phenyl 4-[(2,2-dicyclohexylacetyl)amino]-1-piperidinecarboxylate hydrochloride

To a solution of Intermediate 9 (0.22g) in anhydrous dichloromethane (4mL), under a nitrogen atmosphere, was added Intermediate 58 (0.14g) followed by diisopropylethylamine (0.08mL). After stirring for 4h at 20°C the mixture was diluted with dichloromethane (50mL), washed with saturated aqueous potassium carbonate (3 x 25mL) and 1M hydrochloric acid (40mL), dried over sodium sulphate and evaporated *in vacuo* to give a cream solid. To this was added 4M
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hydrogen chloride in 1,4-dioxane (3mL) and the mixture was stirred for 3h at 20°C. The solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a cream solid (0.24g, 95%). LCMS: R_t 3.05 min; *m/z* 641 (MH⁺).

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Intermediate 11: Tert-butyl (2S)-2-[((2S)-2-[(benzyloxy)carbonyl]amino)-4-methylpentanoyl]amino]-3-(4-hydroxyphenyl)propanoate

To a solution of N-carbobenzyloxy-L-leucine (8.6g) in acetonitrile (150mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (6.83g) and 1-hydroxybenzotriazole (4.81g). After stirring for 30 mins at 20°C L-tyrosine *tert*-butyl ester (7.7g) was added and stirring was continued for 18h. The mixture was concentrated *in vacuo* to ca. 10mL and the residue was partitioned between 1M hydrochloric acid (300mL) and ethyl acetate (150mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (150mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (150mL), water (2 x 150mL) and brine (100mL), dried over sodium sulphate and evaporated *in vacuo*. The residue was co-evaporated with chloroform to give the title compound as a white foam (15g, 96%). LCMS: R_t 3.56 min; *m/z* 485 (MH⁺).

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Intermediate 12: Tert-butyl (2S)-2-[((2S)-2-[(benzyloxy)carbonyl]amino)-4-methyl pentanoyl]amino]-3-(4-[[4-nitrophenoxy]carbonyl]oxy)phenyl)propanoate

To a solution of Intermediate 11 (1.36g) in dichloromethane (15mL), under a nitrogen atmosphere, was added 4-nitrophenyl chloroformate (0.75g) and 4-dimethylaminopyridine (0.47g). The mixture was stirred for 18h at 20°C then diluted with chloroform (50mL), washed with 1M hydrochloric acid (2 x 30mL) and water (30mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting

with petroleum ether/ethyl acetate (4:1 switching to 1:1) to give the title compound as a white solid (1.34g, 74%). LCMS: R_t 3.89 min; m/z 650 (MH^+).

Intermediate 13: 4-[(2S)-2-[((2S)-2-{[(Benzylxy)carbonyl]amino}-4-methyl

5 pentanoyl)amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-morpholinecarboxylate

To a solution of Intermediate 12 (0.34g) in dichloromethane (8mL), under a nitrogen atmosphere, was added morpholine (0.06mL) and diisopropylethylamine (0.15mL). The mixture was stirred for 18h at 20°C then diluted with chloroform (30mL), washed with saturated aqueous potassium carbonate (3 x 40mL), 2M hydrochloric acid (40mL) and water (30mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate/petroleum ether (3:2) to give the title compound as a colourless gum (0.31g, 99%). LCMS: R_t 3.60 min; m/z 598 (MH^+).

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Intermediate 13 (Alternative procedure): 4-[(2S)-2-[((2S)-2-{[(Benzylxy)carbonyl]amino}-4-methylpentanoyl)amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-morpholinecarboxylate

To a solution of triphosgene (2.24g) in anhydrous dichloromethane (50mL), under a nitrogen atmosphere, was added a solution of Intermediate 11 (10g) in anhydrous THF (50mL) followed by diisopropylethylamine (3.94mL). After stirring for 4h at 20°C morpholine (2mL) was added followed by diisopropylethylamine (3.94mL). Stirring was continued for 18h then the mixture was partitioned between 1M hydrochloric acid (100mL) and ethyl acetate (75mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (75mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (50mL), water (50mL) and brine (30mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel

eluting with cyclohexane/ethyl acetate (3:1 switching to 1:1) to give the title compound as a white solid (6.8g, 58%).

5 Intermediate 14: 4-[(2S)-2-[((2S)-2-[(Benzyl oxy)carbonyl]amino)-4-methyl pentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-(aminocarbonyl)-1-piperidinecarboxylate

This was similarly prepared from Intermediate 11 (9g) and isonipecotamide (5.2g). The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate to give the title compound as a white solid (3.52g, 10 30%).

15 Intermediate 14: (Alternative Procedure) 4-[(2S)-2-[(Benzyl oxy) carbonyl]amino]-4-methyl pentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-(aminocarbonyl)-1-piperidinecarboxylate

To a solution of Intermediate 12 (1g) in dichloromethane (20mL), under a nitrogen atmosphere, was added isonipecotamide (0.23g) and diisopropylethylamine (0.43mL). The mixture was stirred for 18h at 20°C then diluted with chloroform (80mL), washed with saturated aqueous potassium carbonate (3 x 50mL), 2M hydrochloric acid (50mL) and water (50mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with petroleum ether/ethyl acetate (3:2) switching to ethyl acetate/methanol (4:1) to give the title compound as a white solid (0.46g, 47%). LCMS: R_t 3.47 min; *m/z* 639 (MH⁺).

25 Intermediate 15: 4-[(2S)-2-[(2S)-2-Amino-4-methylpentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-morpholinecarboxylate

To 10% palladium on carbon, Degussa type E101 (0.09g), under a nitrogen atmosphere, was added a solution of Intermediate 13 (0.3g) in ethanol (20mL) followed by ammonium formate (0.17g). After stirring for 4h at 20°C the mixture

was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (10mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between dichloromethane (50mL) and 1M sodium hydroxide (15mL). The layers were separated and the organic phase 5 was further washed with 1M sodium hydroxide (15mL) and water (15mL), dried over sodium sulphate and evaporated *in vacuo* to give the title compound as a grey gum (0.1g, 41%). LCMS: R_t 2.43 min; *m/z* 464 (MH⁺).

10 Intermediate 16: 4-[(2S)-2-{[(2S)-2-Amino-4-methylpentanoyl]amino}-3-(tert-butoxy)-3-oxopropyl]phenyl 4-(aminocarbonyl)-1-piperidinecarboxylate

This was similarly prepared from Intermediate 14 (0.46g). The title compound was obtained as a pale yellow gum (0.36g, 99%). LCMS: R_t 2.33 min; *m/z* 505 (MH⁺).

15 Intermediate 17: 4-[(2S)-2-{[(2S)-2-{[(Benzyl)oxy]carbonyl}amino]-4-methyl pentanoyl}amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-acetyl-1-piperazine carboxylate

To a solution of triphosgene (0.24g) in anhydrous dichloromethane (5mL), under a nitrogen atmosphere, was added a solution of Intermediate 11 (1g) in anhydrous THF (10mL) followed by diisopropylethylamine (0.43mL). After stirring for 4h at 20°C 1-acetyl piperazine (0.32g) was added followed by diisopropylethylamine (0.43mL). Stirring was continued for 18h then the mixture 20 was partitioned between 1M hydrochloric acid (100mL) and ethyl acetate (75mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (75mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (50mL), water (50mL) and brine (30mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel 25

eluting with ethyl acetate switching to ethyl acetate/ethanol (9:1) to give the title compound as a white foam (1.3g, 99%). LCMS: R_t 3.44 min; m/z 639 (MH^+).

Intermediate 18: 4-[(2S)-2-[((2S)-2-[(Benzylxy)carbonyl]amino)-4-methyl pentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-benzoyl-1-piperazine carboxylate

To a solution of triphosgene (0.24g) in anhydrous dichloromethane (5mL), under a nitrogen atmosphere, was added a solution of Intermediate 11 (1g) in anhydrous THF (10mL) followed by diisopropylethylamine (0.43mL). After stirring for 4h at 20°C Intermediate 56 (0.78g) was added followed by diisopropylethylamine (1.15mL). Stirring was continued for 18h then the mixture was partitioned between 1M hydrochloric acid (100mL) and ethyl acetate (75mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (75mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (50mL), water (50mL) and brine (30mL), dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with ethyl acetate/petroleum ether (1:1 switching to 2:1) to give the title compound as a white foam (1.02g, 71%). LCMS: R_t 3.71 min; m/z 701 (MH^+).

Intermediate 19: 4-[(2S)-2-[((2S)-2-[(Benzylxy)carbonyl]amino)-4-methyl pentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-(1-piperidinylcarbonyl)-1-piperidinecarboxylate

This was similarly prepared from Intermediate 11 (1.81g) and Intermediate 55 (0.91g). The crude product was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol (20:1) to give the title compound as a white foam (1.24g, 47%). LCMS: R_t 3.63 min; m/z 707 (MH^+).

Intermediate 20: 4-[(2S)-2-{[(2S)-2-Amino-4-methylpentanoyl]amino}-3-(tert-butoxy)-3-oxopropyl]phenyl 4-(1-piperidinylcarbonyl)-1-piperidinecarboxylate

To 10% palladium on carbon, Degussa type E101 (0.27g), under a nitrogen atmosphere, was added a solution of Intermediate 19 (1.24g) in ethanol (20mL) followed by ammonium formate (0.77g). After stirring for 4h at 20°C the mixture was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (20mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between dichloromethane (50mL) and 1M sodium hydroxide (15mL). The layers were separated and the organic phase was further washed with 1M sodium hydroxide (15mL) and water (15mL), dried over sodium sulphate and evaporated *in vacuo* to give the title compound as a white foam (0.55g, 54%). LCMS: R_t 2.63 min; *m/z* 573 (MH⁺).

Intermediate 21: 4-[(2S)-2-{[(2S)-2-Amino-4-methylpentanoyl]amino}-3-(tert-butoxy)-3-oxopropyl]phenyl 4-acetyl-1-piperazinecarboxylate hydrochloride

To 10% palladium on carbon, Degussa type E101 (0.4g), under a nitrogen atmosphere, was added a solution of Intermediate 17 (1.28g) in ethanol (30mL) followed by ammonium formate (0.38g). After stirring for 6h at 20°C the mixture was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (20mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between dichloromethane (70mL) and 1M sodium hydroxide (30mL). The layers were separated and the aqueous phase was further extracted with dichloromethane (2 x 50mL). The combined organic extracts were dried over sodium sulphate. The solution was treated with 4M hydrogen chloride in 1,4-dioxane (0.55mL) and evaporated *in vacuo* to give the title compound as a white solid (1.02g, 94%). LCMS: R_t 2.46 min; *m/z* 505 (MH⁺).

Intermediate 22: 4-[(2S)-2-[(2S)-2-Amino-4-methylpentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-benzoyl-1-piperazinecarboxylate hydrochloride

To 10% palladium on carbon, Degussa type E101 (0.3g), under a nitrogen atmosphere, was added a solution of Intermediate 18 (1g) in ethanol (30mL) followed by ammonium formate (0.27g). After stirring for 6h at 20°C the mixture was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (20mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between dichloromethane (70mL) and 1M sodium hydroxide (30mL). The layers were separated and the aqueous phase was further extracted with dichloromethane (2 x 50mL). The combined organic extracts were dried over sodium sulphate. The solution was treated with 4M hydrogen chloride in 1,4-dioxane (0.4mL) and evaporated *in vacuo* to give the title compound as a white solid (0.8g, 100%). LCMS: R_t 2.72 min; *m/z* 567 (MH⁺).

Intermediate 23: 4-[(2S)-2-[(2S)-2-Amino-4-methylpentanoyl]amino]-3-(tert-butoxy)-3-oxopropyl]phenyl 4-morpholinecarboxylate hydrochloride

To 10% palladium on carbon, Degussa type E101 (2.1g), under a nitrogen atmosphere, was added a solution of Intermediate 13 (6.8g) in ethanol (500mL) followed by ammonium formate (4.1g). After stirring for 17h at 20°C the mixture was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (50mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between dichloromethane (150mL) and 1M sodium hydroxide (75mL). The layers were separated and the aqueous phase was further extracted with dichloromethane (2 x 100mL). The combined organic extracts were dried over sodium sulphate. The solution was treated with 1M hydrogen chloride in ether (13mL) and evaporated *in vacuo*. The residue was triturated with ether to give the title compound as a white solid (4.8g, 87%). LCMS: R_t 2.50 min; *m/z* 464 (MH⁺).

Intermediate 24: 4-[(2S)-2-{[(2S)-2-Amino-4-methylpentanoyl]amino}-3-(tert-butoxy)-3-oxopropyl]phenyl 4-(aminocarbonyl)-1-piperidinecarboxylate hydrochloride

To 10% palladium on carbon, Degussa type E101 (1.1g), under a nitrogen atmosphere, was added a solution of Intermediate 14 (3.41g) in ethanol (80mL) followed by ammonium formate (2.1g). After stirring for 3h at 20°C the mixture was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (40mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between chloroform (500mL) and saturated aqueous sodium hydrogen carbonate (200mL). The layers were separated and the aqueous phase was further extracted with chloroform (2 x 100mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (3 x 100mL) and water (2 x 100mL) then dried over sodium sulphate. The solution was treated with 4M hydrogen chloride in 1,4-dioxane (1.5mL) and evaporated *in vacuo*. The residue was azeotroped with toluene (2 x 50mL) to give the title compound as a white solid (2.88g, 100%). LCMS: R_t 2.36 min; *m/z* 505 (MH⁺).

Intermediate 25: Tert-butyl (2S)-2-{[(2S)-2-{2-[2-(tert-butyl)phenoxy]acetyl}amino)-4-methylpentanoyl]amino}-3-(4-hydroxyphenyl)propanoate

To 10% palladium on carbon, Degussa type E101 (0.63g), under a nitrogen atmosphere, was added a solution of Intermediate 11 (2g) in ethanol (20mL) followed by ammonium formate (1.8g). After stirring for 2h at 20°C the mixture was filtered through a pad of Harborlite J2 Filter Aid and the pad was washed with ethanol (50mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between dichloromethane (100mL) and saturated aqueous sodium hydrogen carbonate (50mL). The layers were separated and the organic phase was further washed with saturated aqueous sodium hydrogen carbonate (50mL) and water (50mL), dried over magnesium

sulphate and evaporated *in vacuo* to give a white solid. A solution of this in DMF (5mL) was added to a pre-mixed solution of Intermediate 46 (0.879g), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.809g) and 1-hydroxybenzotriazole (0.578g) in acetonitrile (10mL) which had been stirring under a nitrogen atmosphere for 30 mins at 20°C. Stirring was continued for 18h. The mixture was diluted with ethyl acetate (200mL), washed with 1M hydrochloric acid (3 x 50mL), saturated aqueous sodium hydrogen carbonate (3 x 50mL) and brine (50mL), dried over magnesium sulphate and evaporated *in vacuo* to give the title compound as a white foam (2.1g, 94%). LCMS: R_t 3.83 min; *m/z* 541 (MH⁺).

Intermediate 26: Tert-butyl (2S)-2-{[(2S)-2-({2-[2-(tert-butyl)phenoxy]acetyl} amino)-4-methylpentanoyl]amino}-3-(4-{{(4-nitrophenoxy)carbonyloxy}phenyl} propanoate

To a solution of Intermediate 25 (2.1g) in dichloromethane (20mL), under a nitrogen atmosphere, was added 4-nitrophenyl chloroformate (1.1g) and 4-dimethylaminopyridine (0.69g). The mixture was stirred for 18h at 20°C then diluted with chloroform (80mL), washed with 1M hydrochloric acid (2 x 50mL) and water (50mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with cyclohexane/ethyl acetate (2:1) to give the title compound as a clear oil (2.65g, 97%). LCMS: R_t 4.17 min; *m/z* 706 (MH⁺).

Intermediate 27: 4-[(2S)-2-{[(2S)-2-[(2-Bromoacetyl)amino]-4-methylpentanoyl} amino)-3-(tert-butoxy)-3-oxopropyl]phenyl 4-morpholinecarboxylate

A solution of Intermediate 23 (0.5g) and diisopropylethylamine (0.19mL) in dichloromethane (10mL) was cooled to 0-5°C. To this was added bromoacetyl chloride (0.09mL) followed by diisopropylethylamine (0.19mL) and stirring was continued for 2h. The mixture was diluted with dichloromethane (50mL), washed

with 2M hydrochloric acid (50mL), saturated aqueous sodium hydrogen carbonate (50mL) and brine (30mL), dried over magnesium sulphate and evaporated *in vacuo* to give the title compound as a white foam (0.52g, 89%). LCMS: R_t 3.28 min; *m/z* 584 (MH⁺).

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Intermediate 28: 4-[(2S)-2-((2S)-2-[(2-Bromoacetyl)amino]-4-methylpentanoyl)amino)-3-methoxy-3-oxopropyl]phenyl 4-[(2-phenylacetyl)amino]-1-piperidine carboxylate

To a solution of Intermediate 8 (0.48g) in anhydrous dichloromethane (4mL) was added diisopropylethylamine (0.142mL). The mixture was cooled to 0-5°C and bromoacetyl chloride (0.07mL) was added. Stirring was continued for 1h allowing the reaction to warm to 20°C. The mixture was diluted with dichloromethane (5mL) and washed with saturated aqueous sodium hydrogen carbonate (5mL), water (10mL) and brine (10mL), dried over sodium sulphate and evaporated *in vacuo* to give the title compound as a white solid (0.464g, 85%). LCMS: R_t 3.20 min; *m/z* 672 [M-H]⁻.

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Intermediate 29: (2S)-3-[4-(Allyloxy)phenyl]-2-{{[9H-fluoren-9-ylmethoxy]carbonyl}amino}propanoic acid bound to Wang resin via acid

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To Wang resin (100-200 mesh, 10g) was added a solution of (2S)-3-[4-(allyloxy)phenyl]-2-{{[9H-fluoren-9-ylmethoxy]carbonyl}amino}propanoic acid (8.5g) in DMF (45mL). After 15 mins pyridine (2.4mL) was added followed by 2,6-dichlorobenzoyl chloride (2.75mL). The mixture was shaken for 18h at 20°C. The resin was filtered and washed with DMF (5 x 40mL), dichloromethane (5 x 40mL) and ether (5 x 40mL) then dried *in vacuo*. The amount of (2S)-3-[4-(allyloxy)phenyl]-2-{{[9H-fluoren-9-ylmethoxy]carbonyl}amino} propanoic acid substituted on the resin was calculated to be 0.52 mmol/g.

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Intermediate 30: (2S)-3-[4-(Allyloxy)phenyl]-2-{[(2S)-2-[(9H-fluoren-9-ylmethoxy)carbonyl]amino}-4-methylpentanoyl]amino}propanoic acid bound to Wang resin via acid

Intermediate 29 (2.5mmol) was treated with 20% piperidine in DMF (15mL) and shaken for 1h 30mins at 20°C. The resin was filtered and washed with DMF (5 x 20mL). A solution of Fmoc-leucine (2.8g) in DMF (10mL) was added followed by a solution of benzotriazol-1-yl-oxy-trispyrrolidinophosphonium hexafluoro phosphate (4.1g) in DMF (5mL) and diisopropylethylamine (2.8mL). The mixture was shaken for 18h at 20°C. The resin was filtered and washed with DMF (5 x 20mL), dichloromethane (5 x 20mL) and ether (5 x 20mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was filtered and the filtrate analysed by LCMS: R_t 4.22 min; *m/z* 557 (MH⁺).

15 Intermediate 31: (2S)-3-[4-(Allyloxy)phenyl]-2-{[(2S)-2-[(2-[tert-butyl)phenoxy]acetyl]amino}-4-methylpentanoyl]amino}propanoic acid bound to Wang resin via acid

Intermediate 30 (1mmol) was treated with 20% piperidine in DMF (10mL) and shaken for 1h at 20°C. The resin was filtered and washed with DMF (5 x 10mL). A solution of Intermediate 46 (0.314g) in DMF (10mL) was added followed by a solution of benzotriazol-1-yl-oxy-trispyrrolidinophosphonium hexafluoro phosphate (0.78g) in DMF (5mL) and diisopropylethylamine (0.68mL). The mixture was shaken for 18h at 20°C. The resin was filtered and washed with DMF (5 x 10mL), dichloromethane (5 x 10mL) and ether (5 x 10mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was filtered and the filtrate analysed by LCMS: R_t 4.27 min; *m/z* 525 (MH⁺).

Intermediate 32: (2S)-3-[4-(Allyloxy)phenyl]-2-[((2S)-4-methyl-2-[(2-methylphenoxy)acetyl]amino)pentanoyl]amino]propanoic acid bound to Wang resin via acid

This was similarly prepared from Intermediate 30 (0.97mmol) and (2-methylphenoxy)acetic acid (0.48g). LCMS: R_t 3.89 min; m/z 483 (MH⁺).

Intermediate 33: (2S)-2-{[(2S)-2-{[2-(Tert-butyl)phenoxy]acetyl}amino]-4-methylpentanoyl]amino}-3-(4-[(4-nitrophenoxy)carbonyl]oxy)phenyl)propanoic acid bound to Wang resin via acid

10 Intermediate 31 (1mmol) was treated with a solution of phenylsilane (1mL) in dichloromethane (9mL) followed by tetrakis(triphenylphosphine)palladium(0) (0.1g). The mixture was shaken for 40 mins at 20°C. The resin was filtered and washed with dichloromethane (5 x 10mL) then retreated with a solution of phenylsilane (1mL) in dichloromethane (9mL) followed by

15 tetrakis(triphenylphosphine)palladium(0) (0.1g). After shaking for 40 mins at 20°C the resin was filtered and washed with dichloromethane (5 x 10mL) then treated with a solution of diisopropylethylamine (1.74mL) in 1:1 dichloromethane/THF (16mL). 4-Nitrophenyl chloroformate (2g) was added portionwise and the mixture was shaken for 18h at 20°C. The resin was filtered and washed with dichloromethane (5 x 10mL) and ether (5 x 10mL) then dried *in vacuo*.

20 A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was filtered and the filtrate analysed by LCMS: R_t 4.33 min; m/z 650 (MH⁺).

25 Intermediate 34: (2S)-2-[((2S)-4-Methyl-2-[(2-methylphenoxy)acetyl]amino)pentanoyl]amino]-3-(4-[(4-nitrophenoxy)carbonyl]oxy)phenyl)propanoic acid bound to Wang resin via acid

This was similarly prepared from Intermediate 32 (0.97mmol). LCMS: R_t 3.31 min; m/z 443 (MH⁺).

Intermediate 35: (2S)-2-[((2S)-2-[(9H-Fluoren-9-ylmethoxy)carbonyl]amino)-4-methylpentanoyl]amino]-3-(4-[(4-nitrophenoxy)carbonyl]oxy)phenyl]propanoic acid bound to Wang resin via acid

5 This was similarly prepared from Intermediate 30 (1.05mmol). LCMS: R_t 4.32 min; m/z 682 (MH⁺).

Intermediate 36: (2S)-2-[(2S)-2-[(9H-Fluoren-9-ylmethoxy)carbonyl]amino)-4-methylpentanoyl]amino]-3-[4-([4-(2-furoyl)-1-piperazinyl]carbonyl]oxy)phenyl]propanoic acid bound to Wang resin via acid

10 Intermediate 35 (1.05mmol) was treated with a solution of 1-(2-furoyl)piperazine (0.57g) in 1:1 dichloromethane/THF (9mL) followed by diisopropylethylamine (1.1mL). After shaking for 4h at 20°C the resin was filtered and washed with dichloromethane (5 x 10mL) and ether (5 x 10mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 15 0.5h at 20°C, resin was filtered and the filtrate analysed by LCMS: R_t 3.67 min; m/z 723 (MH⁺).

Intermediate 37: (2S)-3-(4-[(4-[[2-(4-Chlorophenyl)acetyl]amino]-1-piperidinyl]carbonyl]oxy)phenyl)-2-[(2S)-2-[(9H-fluoren-9-ylmethoxy)carbonyl]amino)-4-methylpentanoyl]amino]propanoic acid bound to Wang resin via acid

20 This was similarly prepared from Intermediate 35 (1.7mmol) and Intermediate 53 (1.02g). LCMS: R_t 4.03 min; m/z 795 (MH⁺).

Intermediate 38: (2S)-2-((2S)-2-[(2-Bromoacetyl)amino]-4-methylpentanoyl)amino)-3-[4-([4-(2-furoyl)-1-piperazinyl]carbonyl]oxy)phenyl]propanoic acid bound to Wang resin via acid

25 Intermediate 36 (1.05mmol) was treated with 20% piperidine in DMF (8mL) and shaken for 1h 30mins at 20°C. The resin was filtered and washed with DMF (5 x 10mL). A solution of bromoacetic acid (0.44g) in DMF (8mL) was added follow d

by 1,3-diisopropylcarbodiimide (0.49mL). The mixture was shaken for 18h at 20°C. The resin was filtered and washed with DMF (5 x 10mL), dichloromethane (5 x 10mL) and ether (5 x 10mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was 5 filtered and the filtrate analysed by LCMS: R_t 3.11 min; *m/z* 621 (MH⁺).

Intermediate 39: (2S)-2-((2S)-2-[(2-Bromoacetyl)amino]-4-methylpentanoyl)amino)-3-(4-[(4-[[2-(4-chlorophenyl)acetyl]amino]-1-piperidinyl)carbonyl]oxy)phenyl)propanoic acid bound to Wang resin via acid

This was similarly prepared from Intermediate 37 (0.73mmol). LCMS: R_t 3.43 min; *m/z* 695 (MH⁺).

Intermediate 40: (2S)-3-[4-(Allyloxy)phenyl]-2-((2S)-2-[(2-bromoacetyl)amino]-4-methylpentanoyl)amino)propanoic acid bound to Wang resin via acid

15 Intermediate 30 (0.55mmol) was treated with 20% piperidine in DMF (6mL) and shaken for 1h at 20°C. The resin was filtered and washed with DMF (5 x 10mL). A solution of bromoacetic acid (0.23g) in DMF (3mL) was added followed by 1,3-diisopropylcarbodiimide (0.26mL). The mixture was shaken for 18h at 20°C. The resin was filtered and washed with DMF (5 x 10mL), dichloromethane (5 x 10mL) and ether (5 x 10mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was 20 filtered and the filtrate analysed by LCMS: R_t 3.47 min; *m/z* 455 (MH⁺).

Intermediate 41: (2S)-3-[4-(Allyloxy)phenyl]-2-[(2S)-2-[(2-cyclohexylphenoxy)acetyl]amino]-4-methylpentanoyl)amino]propanoic acid bound to Wang resin via acid

Intermediate 40 (0.55mmol) was treated with DMF (4mL). 2-Cyclohexylphenol (0.97g), potassium carbonate (0.76g) and sodium iodide (0.82g) were added and the mixture was shaken for 40h at 20°C. The resin was filtered and washed

with water (3 x 5mL), DMF (5 x 5mL), dichloromethane (5 x 5mL) and ether (5 x 5mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was filtered and the filtrate analysed by LCMS: R_t 4.49 min; *m/z* 551 (MH⁺).

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Intermediate 42: (2S)-2-[((2S)-2-{[2-(2-Cyclohexylphenoxy)acetyl]amino}-4-methylpentanoyl)amino]-3-(4-{[(4-nitrophenoxy)carbonyl]oxy}phenyl)propanoic acid bound to Wang resin via acid

Intermediate 41 (0.55mmol) was treated with a solution of phenylsilane (1.35mL) in dichloromethane (10mL) followed by tetrakis(triphenylphosphine)palladium(0) (0.063g). The mixture was shaken for 40 mins at 20°C. The resin was filtered and washed with dichloromethane (5 x 10mL) then retreated with a solution of phenylsilane (1.35mL) in dichloromethane (10mL) followed by tetrakis(triphenylphosphine)palladium(0) (0.063g). After shaking for 40 mins at 20°C the resin was filtered and washed with dichloromethane (5 x 10mL) then treated with a solution of diisopropylethylamine (1.9mL) in 1:1 dichloromethane/THF (8mL). 4-Nitrophenyl chloroformate (2.2g) was added portionwise and the mixture was shaken for 18h at 20°C. The resin was filtered and washed with dichloromethane (5 x 10mL) and ether (5 x 10mL) then dried *in vacuo*. A 5mg sample was treated with trifluoroacetic acid/ dichloromethane (1:1) (1mL) for 0.5h at 20°C, resin was filtered and the filtrate analysed by LCMS: R_t 4.54 min; *m/z* 676 (MH⁺).

Intermediate 43: (2-Iodophenoxy)acetic acid

tert-Butyl bromoacetate (4.0mL) was added to a suspension containing 2-iodophenol (4.98g) and potassium carbonate (6.3g) in DMF (40mL). The mixture was stirred for 1h at 20°C under a nitrogen atmosphere and was then partitioned between ethyl acetate (150mL) and water (100mL). The aqueous layer was extracted with fresh ethyl acetate (2 x 80mL) and the combined organic extracts

washed with brine (100mL), dried over magnesium sulphate and evaporated *in vacuo* to give a clear liquid (7.56g). This was dissolved in dichloromethane (20mL) and trifluoroacetic acid (8mL) and the solution stirred for 2h at 20°C. Solvent was evaporated *in vacuo* and the residue triturated in a mixture of cyclohexane/ethyl acetate (5:1) to give the title compound as a white solid (5.19g, 82%). LCMS: R_t 3.02 min; *m/z* 277 [M-H]⁻.

Intermediate 44: {[3-(1-Piperidinylcarbonyl)-2-naphthyl]oxy}acetic acid

This was similarly prepared from 3-(1-piperidinylcarbonyl)-2-naphthol (Griffiths and Hawkins, 1977)((4.98g). The intermediate ester was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) and the title compound was isolated as a white solid (3.2g, 53%). LCMS: R_t 3.74 min; *m/z* 314 (MH⁺).

Intermediate 45: Dibenzo[b,d]furan-4-carboxylic acid

A solution of 1.6M *n*-butyllithium in hexane (18.5mL) was added dropwise to a stirred solution of dibenzofuran (5.0g) in anhydrous THF (25mL) at -78°C under a nitrogen atmosphere. The resulting suspension was allowed to warm to 20°C where it was stirred for 3h. It was then cooled to -78°C and added to a mixture of excess solid carbon dioxide in diethyl ether (250mL) under a nitrogen atmosphere. The resulting white suspension was allowed to stand for 1h at 20°C and was then diluted with 2M sodium hydroxide (500mL). The aqueous extract was washed with ether (3 x 200mL), acidified to pH 1 with 6M hydrochloric acid and extracted with ethyl acetate (3 x 200mL). The combined organic extracts were washed with brine (50mL), dried over magnesium sulphate and evaporated *in vacuo* to give the title compound as a white solid (3.64g, 58%). LCMS: R_t 5.06 min; *m/z* 213 (MH⁺).

Intermediate 46: [2-(Tert-butyl)phenoxy]acetic acid

Methyl bromoacetate (3.0mL) was added to a suspension containing 2-*tert*-butylphenol (5.0mL) and potassium carbonate (10.6g) in DMF (250mL). The mixture was stirred for 20h at 20°C under a nitrogen atmosphere and was then evaporated *in vacuo* to a slurry which was partitioned between ether (200mL) and 1M hydrochloric acid (100mL). The aqueous layer was extracted with more ether (100mL) and the combined organic extracts washed with brine (100mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:9) to give a clear liquid (6.64g). This was dissolved in methanol (100mL) and 2M sodium hydroxide (100mL) and the solution was stirred for 0.5h at 20°C. The methanol was evaporated *in vacuo* and the aqueous residue was washed with diethyl ether (50mL), acidified to pH 1 with 6M hydrochloric acid and extracted with ethyl acetate (2 x 200mL). The combined organic extracts were washed with brine (50mL), dried over magnesium sulphate and evaporated *in vacuo* to give the title compound as a white crystalline mass (5.86g, 95%). LCMS: R_t 3.78 min; *m/z* 207 [M-H]⁻.

Intermediate 47: 4-(2-Methoxy-2-oxoethoxy)benzoic acid

Methyl bromoacetate (1.6mL) was added to a suspension containing *tert*-butyl 4-hydroxybenzoate (Shah *et al.*, 1992) (3.03g), sodium iodide (2.55g) and potassium carbonate (4.2g) in acetonitrile (60mL). The mixture was stirred for 17h at 90°C under a nitrogen atmosphere and then allowed to cool to 20°C. It was then partitioned between water (50mL) and ethyl acetate (100mL) and the organic extract washed with water (2 x 80mL) and brine (60mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with a gradient of ethyl acetate/petroleum ether (1:9) to ethyl acetate/petroleum ether (1:2) to give a pale red gum (3.85g). This was dissolved in dichloromethane (50mL) and trifluoroacetic acid (15mL) was added and the solution was stirred for 3h at

20°C. Solvents were evaporated *in vacuo* to give the title compound as a white solid (2.97g, 91%). LCMS: R_t 2.45 min; *m/z* 211 (MH⁺).

Intermediate 48: [4-(1-Piperidinylcarbonyl)phenoxy]acetic acid

5 To a suspension of Intermediate 47 (2.95g) in acetonitrile (55mL) was added diisopropylethylamine (3.5mL) followed by (1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate (4.5g). The resulting solution was stirred for 10mins at 20°C under a nitrogen atmosphere and then piperidine (1.4mL) was added and the mixture was stirred for 18h at 20°C under a nitrogen atmosphere and then evaporated *in vacuo*. The residue was partitioned between ethyl acetate (100mL) and 8% aqueous sodium hydrogen carbonate (65mL) and the organic extract was washed with 2M hydrochloric acid (50mL) and brine (100mL), dried over magnesium sulphate and evaporated *in vacuo* to give an orange oil (4.05g). This was dissolved in methanol (100mL) and 1M sodium hydroxide (30mL) was added and the mixture stirred for 3h at 20°C. It was then acidified to pH 1 with 1M hydrochloric acid and cooled to 5°C and the precipitate collected by filtration and dried *in vacuo* to give the title compound as a white solid (3.03g, 80%). LCMS: R_t 4.17 min; *m/z* 264 (MH⁺).

20 Intermediate 49: (2-Benzoylphenoxy)acetic acid

Methyl bromoacetate (3.0mL) was added to a suspension containing 2-hydroxybenzophenone (2.3g), potassium carbonate (3.2g) and sodium iodide (2.33g) in acetonitrile (35mL). The mixture was stirred for 18h at 90°C under a nitrogen atmosphere and was then allowed to cool to 20°C. It was then partitioned between ethyl acetate (80mL) and water (60mL) and the organic extract washed with water (2 x 60mL) and brine (60mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate/petroleum ether (1:1) to give a pale yellow oil (3.05g). This was dissolved in methanol (100mL)

and 1M sodium hydroxide (35mL) and the solution was stirred for 18h at 20°C. The solution was acidified to pH 1 with 2M hydrochloric acid and extracted with ethyl acetate (2 x 80mL). The combined organic extracts were washed with water (2 x 70mL), dried over magnesium sulphate and evaporated *in vacuo*. The 5 crude product was purified by flash column chromatography on silica gel eluting with a gradient of ethyl acetate/petroleum ether (1:1) to ethyl acetate/methanol (4:1) to give the title compound as a pale yellow gum (1.62g, 57%). LCMS: R_t 3.41 min; m/z 257 (MH^+).

10 Intermediate 50: [(1-Bromo-2-naphthyl)oxy]acetic acid

This was similarly prepared from 1-bromo-2-naphthol (10.55g). The intermediate ester was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:3) and the title compound was isolated as a pale brown solid (11.36g, 89%). LCMS: R_t 4.17 min; m/z 281 [$M-H^-$].

15

Intermediate 51: [4-(Aminocarbonyl)phenoxy]acetic acid

A solution of 4-formylphenoxyacetic acid (1.86g) and hydroxylamine hydrochloride (1.07g) in 98% formic acid (50mL) was stirred under reflux for 2h and then cooled in an ice bath. The precipitate was collected by filtration, 20 washed with water and dried *in vacuo* to give a white solid (1.1g). A mixture of this with powdered potassium hydroxide (2.3g) in *tert*-butanol (50mL) was stirred under reflux under a nitrogen atmosphere for 4h and then allowed to cool. The mixture was diluted with water (100mL), washed with ethyl acetate (50mL) and acidified to pH 2 with 6M hydrochloric acid. The precipitate was collected by 25 filtration, washed with water and dried *in vacuo* to give the title compound as a white solid (1.06g, 53%). LCMS: R_t 1.90 min; m/z 196 (MH^+).

Intermediate 52: Tert-butyl 4-amino-1-piperidinecarboxylate

Sodium triacetoxyborohydride (30.2g) was added portionwise over 10min to an ice-cooled mixture of 1-(*tert*-butoxycarbonyl)-4-piperidone (20.07g), dibenzylamine (19.7g) and acetic acid (5mL) in dichloromethane (500mL) and stirring was then continued for 16h at 20°C. The solution was then treated
5 cautiously with 2M sodium hydroxide (400mL) and the separated organic layer, dried over magnesium sulphate and evaporated *in vacuo*. The residue was triturated in hexane/ether (2:1) (250mL) to give a white solid (18.75g). This was dissolved in a mixture of THF (50mL), ethanol (50mL) and 2M hydrochloric acid
10 (8mL) and the solution added to a suspension of 20% palladium hydroxide on carbon (5.0g) in ethanol (100mL). The mixture was hydrogenated at 20°C and 1 atmosphere for 17h and was then filtered through a pad of Harborlite J2 Filter Aid and the pad washed with ethanol (100mL). The combined filtrate and washings were evaporated *in vacuo* and the residue dissolved in water (50mL) and adjusted to pH 9 with 2M sodium hydroxide and evaporated *in vacuo*. The
15 residue was leached into a mixture of ethanol (30mL) and chloroform (70mL) and insoluble material removed by filtration. The mother liquors were evaporated *in vacuo* to give the title compound as a colourless oil (10.04g, 49%). LCMS: R_t 1.81 min; *m/z* 201 (MH⁺).

20 Intermediate 53: 2-(4-Chlorophenyl)-N-(4-piperidinyl)acetamide hydrochloride
To a solution of 4-chlorophenylacetic acid (2.55g) in acetonitrile (100mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (3.16g) and 1-hydroxybenzotriazole (2.22g). After stirring for 10 mins at 20°C a solution of Intermediate 52 (3g) in acetonitrile
25 (20mL) was added, and stirring was continued for 18h. The mixture was evaporated *in vacuo* and the residue partitioned between water (100mL) and ethyl acetate (100mL). The organic phase was washed with saturated aqueous sodium hydrogen carbonate (2 x 80mL) and water (50mL), dried over magnesium sulphate and evaporated *in vacuo* to give a pale yellow solid. This

was triturated with ether to give a white solid (4.15g). A portion of this (2.36g) was dissolved in 1,4-dioxane (100mL) and 4M hydrogen chloride in 1,4-dioxane (12mL) was added. The solution was stirred for 18h at 20°C and then a further portion of 4M hydrogen chloride in 1,4-dioxane (8mL) was added. Stirring was continued for a further 18h at 20°C and the solution was evaporated *in vacuo* to give a white solid. This was triturated in ether to give the title compound as a white solid (1.9g, 77%). LCMS: R_t 1.89 min; m/z 253 (MH^+).

Intermediate 54: N-(4-Fluorobenzyl)-4-piperidinecarboxamide hydrochloride

To a solution of 1-*tert*-butoxycarbonylpiperidine-4-carboxylic acid (3.61g) in acetonitrile (25ml), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (3.21g) and 1-hydroxybenzotriazole (2.29g). After stirring for 20 mins at 20°C 4-fluorobenzylamine (2.0mL) was added and stirring was continued for 3h. The mixture was concentrated *in vacuo* and the residue was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (200mL). The layers were separated and the organic phase was washed with 1M hydrochloric acid (3 x 50mL), saturated aqueous sodium hydrogen carbonate (3 x 50mL) and brine (50mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with a gradient of cyclohexane/ethyl acetate (1:1) to neat ethyl acetate to give colourless crystals (5.02g). A portion of this (4.96g) was dissolved in 1,4-dioxane (20mL) and 4M hydrogen chloride in 1,4-dioxane (15mL) was added. The mixture was stirred for 2h at 20°C and the precipitate was collected by filtration, washed with 1,4-dioxane and diethyl ether and dried *in vacuo* to give the title compound as a white hygroscopic solid (3.54g, 83%). LCMS: R_t 1.52 min; m/z 237 (MH^+).

Intermediate 55: 1-(4-Piperidinylcarbonyl)piperidine hydrochloride

This was similarly prepared from 1-*tert*-butoxycarbonylpiperidine-4-carboxylic acid (3.68g) and piperidine (1.6mL). The intermediate amide was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol (10:1) and the title compound was isolated as a white solid (3.26g, 93%). MS: *m/z* 197 (MH^+), TLC: R_f 0.1 [dichloromethane/ethanol/880 ammonia (50:8:1) visualisation with iodoplatinic acid].

Intermediate 56: 1-Benzoylpiperazine

This was similarly prepared from benzoic acid (5.02g) and 1-(*tert*-butoxycarbonyl)piperazine (7.66g) and the title compound was isolated as a white solid (7.7g, 82%). LCMS: R_t 0.51 min; *m/z* 191(MH^+).

Intermediate 57: 2-Cyclohexyl-N-(4-piperidinyl)acetamide

A solution of 4-amino-1-benzylpiperidine (5.0mL), cyclohexaneacetic acid (3.79g) and (1*H*-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate (8.35g) in acetonitrile (60mL) was stirred for 18h at 20°C under a nitrogen atmosphere and was then evaporated *in vacuo* to a syrup. This was partitioned between ethyl acetate (200mL) and saturated aqueous sodium hydrogen carbonate (200mL). The organic extract was washed with saturated aqueous sodium hydrogen carbonate (2 x 100mL) and brine (100mL), dried over magnesium sulphate and evaporated *in vacuo* to give an off-white solid. This was crystallised from cyclohexane to give cream crystals (6.24g). A portion of this (3.8g) was dissolved in ethanol (100mL) and treated with 10% palladium on carbon, Degussa type E101 (1.2g) and ammonium formate (2.24g). The mixture was stirred for 2.5h at 20°C under a nitrogen atmosphere and was then filtered through a pad of Harborlite J2 Filter Aid and the pad washed with ethanol (100mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between chloroform (100mL) and 0.5M potassium

hydroxide (10mL). The layers were separated and the aqueous phase extracted with fresh chloroform (2 x 100mL) and the combined organic extracts dried over magnesium sulphate and evaporated *in vacuo* to give a white solid. This was triturated with ether to give the title compound as a white solid (2.01g, 60%).

5 LCMS: R_t 1.93 min; *m/z* 225 (MH⁺).

Intermediate 58: 2,2-Dicyclohexyl-N-(4-piperidinyl)acetamide

A solution containing dicyclohexylacetic acid (4.75g), diisopropylethylamine (7.5mL) and benzotriazol-1-yl-oxy-trispyrrolidinophosphonium hexafluoro phosphate (11g) in DMF (250mL) was stirred for 10min at 20°C and then 4-amino-1-benzylpiperidine (4.3mL) was added dropwise over 10min. The mixture was stirred for 18h at 20°C and was then diluted with ethyl acetate (200mL) and the precipitate collected by filtration, washed with ethyl acetate (60mL) and water (50mL) and dried *in vacuo* to give a white solid (5.91g). A portion of this (3g) was suspended in ethanol (300mL) and treated with 10% palladium on carbon, Degussa type E101 (1.2g) and ammonium formate (2.68g). The mixture was stirred for 4h at 20°C under a nitrogen atmosphere and was then filtered through a pad of Harborlite J2 Filter Aid and the pad washed with ethanol (50mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between chloroform (200mL) and 0.5M sodium hydroxide (150mL). The layers were separated and the aqueous phase extracted with fresh chloroform (100mL) and the combined organic extracts dried over magnesium sulphate and evaporated *in vacuo* to give a white solid. This was triturated with ice-cold ether to give the title compound as a white solid (1.8, 78%). LCMS: R_t 2.69 min; *m/z* 307 (MH⁺).

Intermediate 59: 2-Phenyl-N-(4-piperidinyl)acetamide

To a solution of phenylacetic acid (3.4g) in acetonitrile (100mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide

hydrochloride (5.28g) and 1-hydroxybenzotriazole (3.72g). After stirring for 30 mins at 20°C 4-amino-1-benzylpiperidine (5.1mL) was added and stirring was continued for 18h. The mixture was concentrated *in vacuo* and the residue was partitioned between 2M hydrochloric acid (100mL) and ethyl acetate (75mL).

5 The layers were separated and the aqueous phase was washed with more ethyl acetate (75mL), basified with solid potassium carbonate and extracted with dichloromethane (2 x 100mL). The combined organic extracts were washed with water (2 x 100mL) and brine (50mL), dried over sodium sulphate and evaporated *in vacuo* to give a white solid (4.8g). A portion of this (4.7g) was dissolved in ethanol (150mL) and treated with 10% palladium on carbon, Degussa type E101 (1.5g) and ammonium formate (2.88g). The mixture was stirred for 4h at 20°C under a nitrogen atmosphere and was then filtered through a pad of Harborlite J2 Filter Aid and the pad washed with ethanol (150mL). The combined filtrate and washings were evaporated *in vacuo* and the residue was partitioned between chloroform (100mL) and 0.5M sodium hydroxide (50mL).

15 The layers were separated and the aqueous phase extracted with fresh chloroform (2 x 100mL) and the combined organic extracts dried over sodium sulphate and evaporated *in vacuo* to give the title compound as a white solid (2.4g, 45%). MS: *m/z* 219 (MH⁺), TLC: R_f 0.16 [dichloromethane/methanol/880 ammonia (40:10:1) visualisation with iodine].

20

Examples

Example 1: (2S)-2-[(2S)-2-{[2-(2-Benzoylphenoxy)acetyl]amino}-4-methyl pentanoyl]amino]-3-{4-[{4-[(2-phenylacetyl)amino]-1-piperidinyl}carbonyl]oxy}phenylpropanoic acid

25 To a solution of 2-hydroxybenzophenone (0.134g) in anhydrous DMF (0.5mL) was added anhydrous potassium carbonate (0.093g) followed by Intermediate 28 (0.152g) and sodium iodide (0.1g). After stirring for 18h at 20°C the mixture was partitioned between saturated aqueous sodium hydrogen carbonate (10mL)

and ethyl acetate (10mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (3 x 10mL). The combined organic extracts were washed with water (20mL) and brine (20mL), dried over sodium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol (10:1) to give a pale yellow solid. To a solution of this in methanol (0.5mL) was added 1M sodium hydroxide (0.22mL). After stirring for 1.5h at 20°C the mixture was partitioned between 2M hydrochloric acid (5mL) and dichloromethane (10mL). The layers were separated and the aqueous phase was further extracted with dichloromethane (2 x 10mL). The combined organic extracts were washed with water (20mL) and brine (20mL), dried over sodium sulphate and evaporated *in vacuo* to give the title compound as a pale yellow foam (0.123g, 73%). LCMS: R_t 3.84 min; *m/z* 775 [M-H]⁻.

Example 2: (2S)-2-((2S)-4-Methyl-2-[(2-[(3-(1-piperidinylcarbonyl)-2-naphthyl]oxy)acetyl]amino]pentanoyl)amino)-3-{4-[(4-[(2-phenylacetyl)amino]-1-piperidinyl)carbonyl]oxy}phenyl]propanoic acid

To a solution of triphosgene (0.04g) in anhydrous dichloromethane (1mL), under a nitrogen atmosphere, was added a solution of Intermediate 3 (0.2g) in anhydrous THF (2mL) followed by diisopropylethylamine (0.07mL). After stirring for 3h at 20°C Intermediate 59 (0.09g) was added followed by diisopropylethylamine (0.07mL). Stirring was continued for 18h then the mixture was partitioned between 2M hydrochloric acid (30mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (20mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (20mL), water (20mL) and brine (20mL), dried over sodium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate switching to ethyl acetate/ethanol (9:1) to give a white foam

(0.19g). To a solution of this (0.15g) in methanol (2mL) was added 2M sodium hydroxide (0.18mL). After stirring for 1h at 20°C the mixture was partitioned between 2M hydrochloric acid (40mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:1) to give the title compound as a white solid (0.12g, 54% from Intermediate 3). LCMS: R_t 3.73 min; *m/z* 834 (MH⁺).

Example 3: (2S)-3-{4-[{(4-[(2,2-Dicyclohexylacetyl)amino]-1-piperidinyl}carbonyl]oxy]phenyl}-2-[(2S)-4-methyl-2-(2-[4-(1-piperidinylcarbonyl)phenoxy]acetyl)amino]pentanoylamino}propanoic acid

To a solution of Intermediate 48 (0.05g) in anhydrous DMF (3mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.04g) and 1-hydroxybenzotriazole (0.03g). After stirring for 30 mins at 20°C Intermediate 10 (0.13g) was added followed by diisopropylethylamine (0.08mL), and stirring was continued for 18h. The mixture was partitioned between 2M hydrochloric acid (40mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (30mL), water (2 x 30mL) and brine (20mL), dried over sodium sulphate and evaporated *in vacuo* to give a cream coloured solid (0.16g). To a solution of this (0.15g) in methanol (2mL) was added 2M sodium hydroxide (0.18mL). After stirring for 1h at 20°C the mixture was partitioned between 2M hydrochloric acid (40mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were dried over sodium sulphate and evaporated *in vacuo*. The crude product was purified

by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:1) to give the title compound as a white solid (0.12g, 62% from Intermediate 10). LCMS: R_t 4.26 min; m/z 872 (MH^+).

5 Example 4: (2S)-2-[(2S)-4-Methyl-2-(2-[4-(1-piperidinylcarbonyl)phenoxy]acetyl)amino]pentanoyl]amino}-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

To a solution of Intermediate 48 (0.06g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.06g) and 1-hydroxybenzotriazole (0.04g). After stirring for 30 mins at 20°C Intermediate 15 (0.1g) was added, and stirring was continued for 18h. The mixture was partitioned between water (20mL) and ethyl acetate (25mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (20mL). The combined organic extracts were washed with water (20mL) and brine (20mL), dried over sodium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with dichloromethane/ethanol/880 ammonia (250:8:1) to give a white sticky solid (0.1g). To this was added trifluoroacetic acid (3mL) and water (3 drops). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.06g, 50%). LCMS: R_t 3.21 min; m/z 653 (MH^+).

20 Example 5: (2S)-3-[4-((4-Aminocarbonyl)-1-piperidinyl]carbonyl)oxy]phenyl]-2-[(2S)-4-methyl-2-(2-[4-(1-piperidinylcarbonyl)phenoxy]acetyl)amino]pentanoyl]amino}propanoic acid

25 This was similarly prepared from Intermediate 48 (0.06g) and Intermediate 16 (0.12g). The crude intermediate ester was purified by flash column chromatography on silica gel eluting with dichloromethane/ethanol/880 ammonia

(500:8:1 switching via 250:8:1 to 100:8:1). The title compound was obtained as a white solid (0.09g, 59%). LCMS: R_t 2.84 min; m/z 694 (MH^+).

Example 6: (2S)-3-[4-({[4-(Aminocarbonyl)-1-piperidiny]carbonyl}oxy)phenyl]-2-[(2S)-2-[(2-(2-benzoylphenoxy)acetyl]amino}-4-methylpentanoyl]amino]propanoic acid

This was similarly prepared from Intermediate 49 (0.07g) and Intermediate 16 (0.11g). The crude intermediate ester was purified by flash column chromatography on silica gel eluting with dichloromethane/ethanol/880 ammonia (500:8:1 switching via 250:8:1 to 100:8:1). The title compound was obtained as a white solid (0.08g, 42%). LCMS: R_t 3.16 min; m/z 687 (MH^+).

Example 7: (2S)-2-[(2S)-2-({2-[4-(Aminocarbonyl)phenoxy]acetyl}amino)-4-methylpentanoyl]amino]-3-[4-({[4-(aminocarbonyl)-1-piperidiny]carbonyl}oxy)phenyl]propanoic acid

This was similarly prepared from Intermediate 51 (0.06g) and Intermediate 16 (0.11g). The crude intermediate ester was purified by flash column chromatography on silica gel eluting with dichloromethane/ethanol/880 ammonia (500:8:1 switching via 250:8:1 and 100:8:1 to 75:8:1). The title compound was obtained as a white solid (0.07g, 55%). LCMS: R_t 2.65 min; m/z 626 (MH^+).

Example 8: (2S)-3-{4-[(4-[(2-Cyclohexylacetyl)amino]-1-piperidiny]carbonyl}oxy]phenyl}-2-[(2S)-2-[(2-(2-iodophenoxy)acetyl]amino}-4-methylpentanoyl]amino]propanoic acid

To a solution of triphosgene (0.058g) in anhydrous dichloromethane (2mL), under a nitrogen atmosphere, was added a solution of Intermediate 4 (0.246g) in anhydrous THF (2mL) followed by diisopropylethylamine (0.11mL). After stirring for 4h at 20°C Intermediate 57 (0.1g) was added followed by diisopropylethylamine (0.07mL). Stirring was continued for 18h then the mixture

was partitioned between 2M hydrochloric acid (50mL) and dichloromethane (50mL). The layers were separated and the organic extract was washed with water (20mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) to give a white foam (0.13g). To a solution of this (0.12g) in methanol (3mL) was added 2M sodium hydroxide (1mL) and water (2mL). After stirring for 18h at 20°C the mixture was partitioned between 2M hydrochloric acid (30mL) and chloroform (30mL). The layers were separated and the organic phase was washed with water (20mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol (4:1) to give the title compound as a white solid (0.064g, 20%). LCMS: R_t 4.12 min; *m/z* 805 (MH⁺).

Example 9: (2S)-3-{4-[{4-[{(2,2-Dicyclohexylacetyl)amino]-1-piperidinyl}carbonyl]oxy]phenyl}-2-[(2S)-2-{[2-(2-iodophenoxy)acetyl]amino}-4-methylpentanoyl]amino]propanoic acid

This was similarly prepared from Intermediate 4 (0.203g) and Intermediate 58 (0.14g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol (9:1) to give the title compound as a white foam (0.153g, 52%). LCMS: R_t 4.45 min; *m/z* 887 (MH⁺).

Example 10: (2S)-2-((2S)-2-[(Dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

To a solution of Intermediate 6 (0.165g) in dichloromethane (5mL), under a nitrogen atmosphere, was added morpholine (0.04mL) and diisopropylethylamine (0.05mL). After stirring for 30 mins at 20°C the solution was diluted with dichloromethane (50mL) and washed with saturated aqueous potassium carbonate (3 x 30mL), 1M hydrochloric acid (2 x 40mL) and water

(30mL), dried over magnesium sulphate and evaporated *in vacuo* to give a white foam (0.143g). To a solution of this (0.14g) in methanol (2mL) was added 1M sodium hydroxide (2mL) and the mixture was stirred for 30 mins at 20°C, then partitioned between 1M hydrochloric acid (40mL) and ethyl acetate (50mL). The
5 organic extract was washed with brine (30mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol (4:1) to give the
title compound as a white solid (0.1g, 69%). LCMS: R_t 3.85 min; m/z 602 (MH⁺).
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Example 11: (2S)-2-((2S)-2-[(Dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methyl pentanoyl)amino)-3-[4-((4-(2-furoyl)-1-piperazinyl)carbonyl)oxy]phenyl propanoic acid

To a solution of Intermediate 6 (0.13g) in dichloromethane (5mL), under a
15 nitrogen atmosphere, was added 1-(2-furoyl)piperazine (0.04g) and diisopropylethylamine (0.04mL). After stirring for 3h at 20°C the solution was diluted with dichloromethane (20mL) and washed with saturated aqueous potassium carbonate (3 x 20mL), 1M hydrochloric acid (2 x 20mL) and water (20mL), dried over magnesium sulphate and evaporated *in vacuo* to give a white
20 foam (0.153g). To a solution of this (0.15g) in methanol (2mL) was added 1M sodium hydroxide (2mL) and the mixture was stirred for 30 mins at 20°C, then partitioned between 1M hydrochloric acid (20mL) and ethyl acetate (20mL). The organic extract was washed with brine (20mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column
25 chromatography on silica gel eluting with chloroform/methanol (4:1) to give the
title compound as a white solid (0.126g, 92%). LCMS: R_t 3.85 min; m/z 695 (MH⁺).

Example 12: (2S)-3-(4-[(4-Benzoyl-1-piperazinyl)carbonyloxy]phenyl)-2-[(2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl]amino)propanoic acid

To a solution of Intermediate 6 (0.172g) in dichloromethane (4mL), under a nitrogen atmosphere, was added Intermediate 56 (0.084g) and diisopropylethylamine (0.2mL). After stirring for 3h at 20°C the solution was diluted with dichloromethane (50mL) and washed with saturated aqueous potassium carbonate (3 x 50mL), 1M hydrochloric acid (2 x 50mL) and water (50mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (4:1) to give a white foam. To a solution of this in methanol (2mL) was added 1M sodium hydroxide (2mL) and the mixture was stirred for 1h at 20°C, then partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (50mL). The organic extract was washed with brine (50mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol (4:1) to give the title compound as a white solid (0.041g, 23%). LCMS: R_t 3.72 min; *m/z* 705 (MH⁺).

Example 13: (2S)-2-[(2S)-2-[(Dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl]amino)-3-{4-[(4-[(2-phenylacetyl)amino]-1-piperidinyl]carbonyloxy]phenyl}propanoic acid

To a solution of Intermediate 45 (0.055g) in acetonitrile (2mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.052g) and 1-hydroxybenzotriazole (0.038g). After stirring for 30 mins at 20°C Intermediate 8 (0.15g) was added followed by diisopropylethylamine (0.047mL), and stirring was continued for 18h. The mixture was diluted with chloroform (100mL) and washed with 1M hydrochloric acid (3 x 50mL), saturated aqueous sodium hydrogen carbonate (3 x 50mL) and

water (50mL), dried over magnesium sulphate and evaporated *in vacuo* to give a white foam (0.189g). To a solution of this (0.176g) in methanol (4mL) was added 1M sodium hydroxide (1mL) and the mixture was stirred for 2h at 20°C; then partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (200mL). The organic extract was washed with brine (30mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with a gradient of chloroform/methanol (9:1) to chloroform/methanol (4:1) to give the title compound as a white solid (0.103g, 79%). LCMS: R_t 4.00 min; *m/z* 733 (MH⁺).

Example 14: (2S)-2-[((2S)-2-{[2-(2-iodophenoxy)acetyl]amino}-4-methyl pentanoyl)amino]-3-{4-[{(4-[(2-phenylacetyl)amino]-1-piperidinyl}carbonyl]oxy] phenyl}propanoic acid

This was similarly prepared from Intermediate 43 (0.073g) and Intermediate 8 (0.15g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol (6:1) to give the title compound as a white solid (0.103g, 53%). LCMS: R_t 3.84 min; *m/z* 799 (MH⁺).

Example 15: (2S)-3-(4-[(4-Acetyl-1-piperazinyl)carbonyl]oxy)phenyl)-2-[((2S)-2- {[2-(2-iodophenoxy)acetyl]amino}-4-methylpentanoyl)amino]propanoic acid

To a solution of Intermediate 43 (0.07g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.05g) and 1-hydroxybenzotriazole (0.04g). After stirring for 30 mins at 20°C Intermediate 21 (0.135g) was added followed by diisopropylethylamine (0.05mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (2 x 50mL),

5 dried over sodium sulphate and evaporated *in vacuo*. The residue was co-evaporated with dichloromethane to give a white foam. To this was added trifluoroacetic acid (2mL) and water (3 drops). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.143g, 83%). LCMS: R_t 3.12 min; *m/z* 709 (MH⁺).

10 Example 16: (2S)-3-(4-[(4-Acetyl-1-piperazinyl)carbonyloxy]phenyl)-2-[(2S)-2-
{{2-[2-(tert-butyl)phenoxy]acetyl}amino}-4-methylpentanoyl]amino]propanoic
acid

15 To a solution of Intermediate 46 (0.052g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.05g) and 1-hydroxybenzotriazole (0.04g). After stirring for 30 mins at 20°C Intermediate 21 (0.135g) was added followed by diisopropylethylamine (0.05mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (2 x 50mL),
20 dried over sodium sulphate and evaporated *in vacuo*. The residue was co-evaporated with dichloromethane to give a white foam. To this was added trifluoroacetic acid (2mL) and water (3 drops). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.115g, 74%). LCMS: R_t 3.31 min; *m/z* 639 (MH⁺).

25 Example 17: (2S)-3-(4-[(4-Acetyl-1-piperazinyl)carbonyloxy]phenyl)-2-[(2S)-4-methyl-2-[(2-(2-methylphenoxy)acetyl]amino]pentanoyl]amino]propanoic acid

To a solution of (2-methylphenoxy)acetic acid (0.042g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.05g) and 1-hydroxybenzotriazole (0.04g). After stirring for 30 mins at 20°C Intermediate 21 (0.135g) was added followed by diisopropylethylamine (0.05mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (2 x 50mL), dried over sodium sulphate and evaporated *in vacuo*. The residue was co-evaporated with dichloromethane to give a white foam. To this was added trifluoroacetic acid (2mL) and water (3 drops). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue triturated with ether to give the title compound as a white solid (0.124g, 86%). LCMS: R_t 3.10 min; m/z 15 597 (MH⁺).

Example 18: (2S)-3-(4-{[(4-Acetyl-1-piperazinyl)carbonyl]oxy}phenyl)-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)propanoic acid

To a solution of Intermediate 45 (0.053g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.05g) and 1-hydroxybenzotriazole (0.04g). After stirring for 30 mins at 20°C Intermediate 21 (0.135g) was added followed by diisopropylethylamine (0.05mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (2 x 50mL), dried over sodium sulphate and evaporated *in vacuo*. The residue was co-

evaporated with dichloromethane to give a white foam. To this was added trifluoroacetic acid (2mL) and water (3 drops). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.127g, 83%). LCMS: R_t 3.33 min; *m/z* 5 643 (M⁺).

Example 19: (2S)-3-(4-{[(4-Benzoyl-1-piperazinyl)carbonyloxy]phenyl}-2-[(2S)-2-[(2-iodophenoxy)acetyl]amino]-4-methylpentanoyl]amino]propanoic acid
This was similarly prepared from Intermediate 43 (0.07g) and Intermediate 22 10 (0.151g). The title compound was obtained as a white solid (0.152g, 81%). LCMS: R_t 3.58 min; *m/z* 771 (M⁺).

Example 20: (2S)-3-(4-{[(4-Benzoyl-1-piperazinyl)carbonyloxy]phenyl}-2-[(2S)-2-[(2-(tert-butyl)phenoxy)acetyl]amino]-4-methylpentanoyl]amino]propanoic 15 acid

To a solution of Intermediate 46 (0.052g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.05g) and 1-hydroxybenzotriazole (0.04g). After stirring for 30 mins at 20°C Intermediate 22 (0.151g) was added followed by 20 diisopropylethylamine (0.05mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (30mL). The layers were separated and the aqueous phase was further extracted with ethyl acetate (30mL). The combined organic extracts were washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (2 x 50mL), 25 dried over sodium sulphate and evaporated *in vacuo*. The residue was co-evaporated with dichloromethane to give a white foam. To this was added trifluoroacetic acid (2mL) and water (3 drops). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to

give the title compound as a white foam (0.17g, 90%). LCMS: R_t 3.61 min; m/z 701 (MH^+).

Example 21: (2S)-3-(4-[(4-Benzoyl-1-piperazinyl)carbonyloxy]phenyl)-2-[(2S)-
5 4-methyl-2-[(2-(2-methylphenoxy)acetyl]amino]pentanoyl]amino]propanoic acid

To a solution of (2-methylphenoxy)acetic acid (0.472g) in acetonitrile (30mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.56g) and 1-hydroxybenzotriazole (0.4g). After stirring for 30 mins at 20°C a solution of Intermediate 22 (1.5g) in acetonitrile (25mL) was added and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (75mL). The layers were separated and the organic phase was washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (50mL), dried over sodium sulphate and evaporated *in vacuo* to give a white foam. To a solution of this in chloroform (12mL) was added trifluoroacetic acid (6mL). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was co-evaporated with chloroform and ether to give the title compound as a white foam (0.17g, 90%). LCMS: R_t 3.44 min; m/z 659 (MH^+).

20 Example 22: (2S)-3-(4-[(4-Benzoyl-1-piperazinyl)carbonyloxy]phenyl)-2-[(2S)-
2-[(2-(2,4-dichlorophenoxy)acetyl]amino]-4-methylpentanoyl]amino]propanoic
acid

This was similarly prepared from 2,4-dichlorophenoxyacetic acid (0.055g) and Intermediate 22 (0.151g). The title compound was obtained by trituration with ether as a white solid (0.129g, 75%). LCMS: R_t 3.52 min; m/z 713 (MH^+).

Example 23: (2S)-2-[(2S)-2-[(2-(2-iodophenoxy)acetyl]amino]-4-methyl
pentanoyl]amino]-3-{4-[(4-morpholinylcarbonyloxy]phenyl}propanoic acid

To a solution of Intermediate 43 (0.556g) in acetonitrile (40mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.383g) and 1-hydroxybenzotriazole (0.27g). After stirring for 30 mins at 20°C Intermediate 23 (1g) was added followed by diisopropylethylamine (0.35mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (75mL). The layers were separated and the organic phase was washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (50mL), dried over sodium sulphate and evaporated *in vacuo* to give a white foam. To a solution of this in dichloromethane (20mL) was added trifluoroacetic acid (20mL) and water (1mL). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (1.15g, 92%). LCMS: R_t 3.68 min; m/z 668 (MH^+).

Example 24: (2S)-2-{[(2S)-2-{[2-(Tert-butyl)phenoxy]acetyl}amino]-4-methyl pentanoyl}amino}-3-{4-[{(4-morpholinylcarbonyl)oxy}phenyl]propanoic acid
To a solution of Intermediate 46 (0.416g) in acetonitrile (40mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.383g) and 1-hydroxybenzotriazole (0.27g). After stirring for 30 mins at 20°C Intermediate 23 (1g) was added followed by diisopropylethylamine (0.35mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (75mL). The layers were separated and the organic phase was washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (50mL), dried over sodium sulphate and evaporated *in vacuo* to give a white foam. To a solution of this in dichloromethane (20mL) was added trifluoroacetic acid (20mL) and water (1mL). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.63g, 53%). LCMS: R_t 3.90 min; m/z 598 (MH^+).

Example 25: (2S)-2-[((2S)-4-Methyl-2-{[2-(2-methylphenoxy)acetyl]amino}pentanoyl)amino]-3-{4-[{(4-morpholinylcarbonyl)oxy}phenyl]propanoic acid}

To a solution of (2-methylphenoxy)acetic acid (0.332g) in acetonitrile (40mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.383g) and 1-hydroxybenzotriazole (0.27g). After stirring for 30 mins at 20°C Intermediate 23 (1g) was added followed by diisopropylethylamine (0.35mL) and stirring was continued for 18h. The mixture was partitioned between 1M hydrochloric acid (50mL) and ethyl acetate (75mL). The layers were separated and the organic phase was washed with saturated aqueous sodium hydrogen carbonate (40mL) and water (50mL), dried over sodium sulphate and evaporated *in vacuo* to give a white foam. To a solution of this in dichloromethane (20mL) was added trifluoroacetic acid (20mL) and water (1mL). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.895g, 80%). LCMS: R_t 3.31 min; *m/z* 556 (MH⁺).

Example 26: (2S)-3-[4-({[4-(Aminocarbonyl)-1-piperidinyl]carbonyl}oxy)phenyl]-2-[(2S)-2-{[2-(2-iodophenoxy)acetyl]amino}-4-methylpentanoyl)amino]propanoic acid

This was similarly prepared from Intermediate 43 (0.06g) and Intermediate 24 (0.1g). The title compound was obtained as a white solid (0.07g, 56%). LCMS: R_t 3.33 min; *m/z* 709 (MH⁺).

Example 27: (2S)-3-[4-({[4-(Aminocarbonyl)-1-piperidinyl]carbonyl}oxy)phenyl]-2-[(2S)-4-methyl-2-{[2-(2-methylphenoxy)acetyl]amino}pentanoyl)amino]propanoic acid

To a solution of (2-methylphenoxy)acetic acid (0.345g) in acetonitrile (50mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.4g) and 1-hydroxybenzotriazole (0.3g). After

stirring for 30 mins at 20°C Intermediate 24 (1g) was added followed by diisopropylethylamine (0.35mL) and stirring was continued for 18h. The mixture was concentrated *in vacuo* and the residue was partitioned between 1M hydrochloric acid (100mL) and ethyl acetate (300mL). The layers were separated and the organic phase was washed with 1M hydrochloric acid (2 x 100mL), saturated aqueous sodium hydrogen carbonate (3 x 100mL) and brine (100mL), dried over magnesium sulphate and evaporated *in vacuo* to give a white solid. To a solution of this in chloroform (5mL) was added trifluoroacetic acid (5mL) and water (1mL). After stirring for 3h at 20°C the solvent was evaporated *in vacuo* and the residue was azeotroped with toluene (2 x 20mL) then triturated with ether to give the title compound as a white solid (1.06g, 96%). LCMS: R_t 3.20 min; *m/z* 597 (MH⁺).

Example 28: (2S)-3-[4-({[4-(Aminocarbonyl)-1-piperidinyl]carbonyl}oxy)phenyl]-2-
15 ((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)
propanoic acid

To a solution of Intermediate 45 (0.438g) in acetonitrile (50mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.4g) and 1-hydroxybenzotriazole (0.29g). After stirring for 30 mins at 20°C Intermediate 24 (1g) was added followed by diisopropylethylamine (0.35mL) and stirring was continued for 18h. The mixture was concentrated *in vacuo* and the residue was partitioned between 1M hydrochloric acid (100mL) and ethyl acetate (300mL). The layers were separated and the organic phase was washed with 1M hydrochloric acid (2 x 100mL), saturated aqueous sodium hydrogen carbonate (3 x 100mL) and brine (100mL), dried over magnesium sulphate and evaporated *in vacuo* to give a white solid. To a solution of this in chloroform (5mL) was added trifluoroacetic acid (5mL) and water (1mL). After stirring for 3h at 20°C the solvent was evaporated *in vacuo* and the residue was

azeotroped with toluene (2 x 20mL) then triturated with ether to give the title compound as a white solid (0.95g, 80%). LCMS: R_t 3.48 min; m/z 643 (MH⁺).

Example 29: (2S)-2-{[(2S)-2-({2-[2-(Tert-butyl)phenoxy]acetyl}amino)-4-methyl pentanoyl]amino}-3-[4-({[4-(1-piperidinylcarbonyl)-1-piperidinyl]carbonyl}oxy) phenyl]propanoic acid

To a solution of Intermediate 46 (0.1g) in acetonitrile (5mL), under a nitrogen atmosphere, was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.09g) and 1-hydroxybenzotriazole (0.063g). After stirring for 30 mins at 20°C Intermediate 20 (0.18g) was added and stirring was continued for 18h. The mixture was partitioned between water (20mL) and ethyl acetate (20mL). The layers were separated and the organic phase was washed with saturated aqueous sodium hydrogen carbonate (2 x 30mL), water (30mL) and brine (30mL), dried over sodium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol (20:1) to give a clear oil. To a solution of this in dichloromethane (8mL) was added trifluoroacetic acid (2mL). After stirring for 2h at 20°C the solvent was evaporated *in vacuo* and the crude product purified by flash column chromatography on silica gel eluting with dichloromethane/methanol/acetic acid/water (240:15:3:2) to give the title compound as a white foam (0.08g, 36%). LCMS: R_t 4.07 min; m/z 707 (MH⁺).

Example 30: (2S)-2-{[(2S)-4-Methyl-2-{[2-(2-methylphenoxy)acetyl]amino} pentanoyl]amino}-3-[4-({[4-(1-piperidinylcarbonyl)-1-piperidinyl]carbonyl}oxy) phenyl]propanoic acid

This was similarly prepared from (2-methylphenoxy)acetic acid (0.09g) and Intermediate 20 (0.3g). The crude product was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol/acetic

acid/water (240:15:3:2) to give the title compound as a white foam (0.116g, 34%). LCMS: R_t 3.56 min; m/z 665 (MH^+).

5 Example 31: (2S)-2-((2S)-2-[(Dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino}-3-[4-((4-(1-piperidinylcarbonyl)-1-piperidinyl)carbonyl)oxy]phenyl]propanoic acid

This was similarly prepared from Intermediate 45 (0.1g) and Intermediate 20 (0.176g). The crude product was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol/acetic acid/water (180:15:3:2) to give the title compound as a white foam (0.075g, 35%). LCMS: R_t 4.09 min; m/z 711 (MH^+).

10 Example 32: (2S)-2-((2S)-2-((2-[(1-Bromo-2-naphthyl)oxy]acetyl)amino)-4-methylpentanoyl)amino}-3-[4-((4-(1-piperidinylcarbonyl)-1-piperidinyl)carbonyl)oxy]phenyl]propanoic acid

15 This was similarly prepared from Intermediate 50 (0.124g) and Intermediate 20 (0.168g). The crude product was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol/acetic acid/water (200:15:3:2) to give the title compound as a white foam (0.055g, 24%). LCMS: R_t 4.19 min; m/z 779 (MH^+).

20 Example 33: (2S)-3-[4-((4-(Aminocarbonyl)-1-piperidinyl)carbonyl)oxy]phenyl]-2-((2S)-2-((2-[2-(tert-butyl)phenoxy]acetyl)amino)-4-methylpentanoyl)amino}propanoic acid

25 To a solution of Intermediate 26 (0.47g) in dichloromethane (8mL), under a nitrogen atmosphere, was added isonipecotamide (0.106g) and diisopropylethylamine (0.2mL). The mixture was stirred for 18h at 20°C then diluted with chloroform (100mL), washed with saturated aqueous potassium carbonate (3 x 50mL), 1M hydrochloric acid (3 x 50mL) and water (50mL), dried

over magnesium sulphate and evaporated *in vacuo* to give a white foam. To a solution of this in chloroform (3mL) was added trifluoroacetic acid (3mL). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.223g, 52%).

5 LCMS: R_t 3.35 min; *m/z* 639 (MH⁺).

Example 34: (2S)-2-{[(2S)-2-((2-[2-(Tert-butyl)phenoxy]acetyl)amino)-4-methyl pentanoyl]amino}-3-{4-[(4-[(4-fluorobenzyl)amino]carbonyl)-1-piperidinyl] carbonyl]oxy}phenyl propanoic acid

0 This was similarly prepared from Intermediate 26 (0.312g) and Intermediate 54 (0.181g). The title compound was obtained as a white solid (0.187g, 57%).

LCMS: R_t 3.71 min; *m/z* 747 (MH⁺).

Example 35: (2S)-2-{[(2S)-2-[(2-(2,4-Dichlorophenoxy)acetyl)amino]-4-methyl pentanoyl]amino}-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

15 To a suspension of anhydrous potassium carbonate (0.057g) and sodium iodide (0.051g) in anhydrous DMF (1mL) was added 2,4-dichlorophenol (0.166g) followed by Intermediate 27 (0.2g). The mixture was stirred for 18h at 20°C then partitioned between saturated aqueous sodium hydrogen carbonate (10mL) and ethyl acetate (10mL). The layers were separated and the organic phase was further washed with saturated aqueous sodium hydrogen carbonate (10mL) and brine (10mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) to give a white foam. To a solution of this in dichloromethane (2mL) was added trifluoroacetic acid (2mL). After stirring for 2h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.146g, 70%). LCMS: R_t 3.70 min; *m/z* 610 (MH⁺).

Example 36: (2S)-2-[((2S)-2-[(2-(2-Benzoylphenoxy)acetyl]amino)-4-methylpentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

This was similarly prepared from 2-hydroxybenzophenone (0.2g) and Intermediate 27 (0.2g). The title compound was obtained as a pale yellow foam (0.057g, 26%). LCMS: R_t 3.60 min; m/z 646 (MH^+).

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Example 37: (2S)-2-[(2S)-4-Methyl-2-[(2-(2-propylphenoxy)acetyl]amino]pentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

This was similarly prepared from 2-propylphenol (0.14mL) and Intermediate 27 (0.2g). The title compound was obtained as a white solid (0.141g, 70%). LCMS: R_t 3.71 min; m/z 584 (MH^+).

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Example 38: (2S)-2-[(2S)-2-[(2-[(1-Bromo-2-naphthyl)oxy]acetyl]amino)-4-methylpentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

This was similarly prepared from 1-bromo-2-naphthol (0.23g) and Intermediate 27 (0.2g). The title compound was obtained as a white solid (0.11g, 48%). LCMS: R_t 3.91 min; m/z 670 (MH^+).

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Example 39: (2S)-2-[(2S)-2-[(2-(2-Cyclohexylphenoxy)acetyl]amino)-4-methylpentanoyl]amino]-3-{4-[(4-morpholinylcarbonyl)oxy]phenyl}propanoic acid

To a suspension of anhydrous potassium carbonate (0.1g) and sodium iodide (0.06g) in anhydrous DMF (1mL) was added 2-cyclohexylphenol (0.12g) followed by Intermediate 27 (0.2g). The mixture was stirred for 18h at 20°C then partitioned between saturated aqueous sodium hydrogen carbonate (10mL) and ethyl acetate (10mL). The layers were separated and the organic phase was further washed with saturated aqueous sodium hydrogen carbonate (10mL) and brine (10mL), dried over magnesium sulphate and evaporated *in vacuo*. The crude material was purified by flash column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) to give a white foam. To a solution of this in

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dichloromethane (3mL) was added trifluoroacetic acid (3mL). After stirring for 2h at 20°C the solvent was evaporated *in vacuo* and the residue was azeotroped with toluene then triturated with ether to give the title compound as a white solid (0.118g, 55%). LCMS: R_t 4.16 min; *m/z* 624 (MH⁺).

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Example 40: (2S)-2-[(2S)-2-{[(Benzyl oxy)carbonyl]amino}-4-methylpentanoyl]amino]-3-[4-[(4-morpholinyl carbonyl)oxy]phenyl]propanoic acid

To a solution of Intermediate 13 (0.19g) in chloroform (2mL) was added trifluoroacetic acid (2mL). After stirring for 4h at 20°C the solvent was evaporated *in vacuo* and the residue was triturated with ether to give the title compound as a white solid (0.156g, 90%). LCMS: R_t 3.22 min; *m/z* 542 (MH⁺).

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Example 41: (2S)-3-[4-({[4-(2-Furoyl)-1-piperazinyl]carbonyl}oxy)phenyl]-2-[(2S)-2-{[2-(2-iodophenoxy)acetyl]amino}-4-methylpentanoyl]amino]propanoic acid

15 Intermediate 38 (0.26mmol) was treated with DMF (4mL). 2-Iodophenol (0.57g), potassium carbonate (0.36g) and sodium iodide (0.39g) were added and the mixture was shaken for 16h at 20°C. The resin was filtered and washed with water (2 x 5mL), DMF (5 x 5mL) and dichloromethane (5 x 5mL) then treated with 1:1 trifluoroacetic acid/ dichloromethane (4mL). After 30 mins the resin was filtered and the filtrate was evaporated *in vacuo*. The residue was azeotroped with toluene (5mL) then triturated with ether. The crude product was crystallised from acetonitrile to give the title compound as a white solid (0.043g).
20 LCMS: R_t 3.50 min; *m/z* 761 (MH⁺).

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25 Example 42: (2S)-2-[(2S)-2-({2-[2-(Tert-butyl)phenoxy]acetyl}amino)-4-methylpentanoyl]amino]-3-[4-({[4-(2-furoyl)-1-piperazinyl]carbonyl}oxy)phenyl]propanoic acid

Intermediate 38 (0.26mmol) was treated with DMF (4mL). 2-*tert*-butyl phenol (0.4mL), potassium carbonate (0.36g) and sodium iodide (0.39g) were added

and the mixture was shaken for 16h at 20°C. The resin was filtered and washed with water (2 x 5mL), DMF (5 x 5mL) and dichloromethane (5 x 5mL) then treated with 1:1 trifluoroacetic acid/ dichloromethane (4mL). After 30 mins the resin was filtered and the filtrate was evaporated *in vacuo*. The residue was
5 azeotroped with toluene (5mL) then triturated with ether. The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:1) to give the title compound as a white solid (0.04g). LCMS: R_t 3.63 min; m/z 691 (MH⁺).

10 Example 43: (2S)-2-[((2S)-2-[(2-Cyclohexylphenoxy)acetyl]amino)-4-methyl pentanoyl]amino]-3-[4-((4-(2-furoyl)-1-piperazinyl)carbonyl)oxy)phenyl] propanoic acid

This was similarly prepared from Intermediate 38 (0.26mmol) and 2-cyclohexyl phenol (0.46g). The crude product was purified using a solid phase extraction 15 cartridge containing reverse phase silica eluting with a chloroform/methanol gradient (increasing from 98:2 to 80:20) to give the title compound as a cream solid (0.037g). LCMS: R_t 3.83 min; m/z 717 (MH⁺).

20 Example 44: (2S)-2-[(2S)-2-((2-[(1-Bromo-2-naphthyl)oxy]acetyl)amino)-4-methylpentanoyl]amino]-3-[4-((4-(2-furoyl)-1-piperazinyl)carbonyl)oxy)phenyl] propanoic acid

This was similarly prepared from Intermediate 38 (0.26mmol) and 1-bromo-2-naphthol (0.58g). The crude product was crystallised from acetonitrile to give the 25 title compound as a cream coloured solid (0.064g). LCMS: R_t 3.69 min; m/z 763 (MH⁺).

Example 45: (2S)-3-(4-((4-[(2-(4-Chlorophenyl)acetyl]amino)-1-piperidinyl) carbonyl)oxy)phenyl)-2-[((2S)-2-[(2-(2-cyclohexylphenoxy)acetyl]amino)-4-methylpentanoyl]amino]propanoic acid

This was similarly prepared from Intermediate 39 (0.29mmol) and 2-cyclohexyl phenol (0.48g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:0.5) to give the title compound as a white solid (0.073g). LCMS: R_t 4.13 min; m/z 789 (MH^+).

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Example 46: (2S)-2-[((2S)-2-{{[2-(2-Benzoylphenoxy)acetyl]amino}-4-methyl pentanoyl)amino]-3-(4-{{[4-({[2-(4-chlorophenyl)acetyl]amino}-1-piperidinyl) carbonyl]oxy}phenyl)propanoic acid}

This was similarly prepared from Intermediate 39 (0.29mmol) and 2-hydroxybenzophenone (0.55g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:0.5) to give the title compound as a white solid (0.065g). LCMS: R_t 3.75 min; m/z 811 (MH^+).

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Example 47: (2S)-3-(4-{{[4-({[2-(4-Chlorophenyl)acetyl]amino}-1-piperidinyl) carbonyl]oxy}phenyl)-2-[((2S)-2-{{[2-(2-iodophenoxy)acetyl]amino}-4-methyl pentanoyl)amino]propanoic acid}

Intermediate 37 (0.27mmol) was treated with 20% piperidine in DMF (5mL) and shaken for 1h at 20°C. The resin was filtered and washed with DMF (5 x 5mL). A solution of Intermediate 43 (0.154g) in DMF (3mL) was added followed by a solution of benzotriazol-1-yl-oxy-trispyrrolidinophosphonium hexafluoro phosphate (0.285g) in DMF (2mL) and diisopropylethylamine (0.26mL). The mixture was shaken for 18h at 20°C. The resin was filtered and washed with DMF (5 x 5mL) and dichloromethane (5 x 5mL), then treated with 1:1 trifluoroacetic acid/ dichloromethane (5mL). After 30 mins the resin was filtered and the filtrate was evaporated *in vacuo*. The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:0.5) to give the title compound as a white solid (0.083g). LCMS: R_t 3.76 min; m/z 833 (MH^+).

Example 48: (2S)-2-[(2S)-2-(2-[2-(Tert-butyl)phenoxy]acetyl)amino]-4-methylpentanoyl]amino}-3-(4-[(4-[(2-(4-chlorophenyl)acetyl)amino]-1-piperidinyl)carbony]oxy}phenyl)propanoic acid

This was similarly prepared from Intermediate 37 (0.27mmol) and Intermediate 46 (0.115g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:0.5) to give the title compound as a white solid (0.107g). LCMS: R_t 3.93 min; m/z 763 (MH^+).

Example 49: (2S)-3-(4-[(4-[(2-(4-Chlorophenyl)acetyl)amino]-1-piperidinyl)carbony]oxy}phenyl)-2-((2S)-2-[(dibenzo[b,d]furan-4-ylcarbonyl)amino]-4-methylpentanoyl)amino)propanoic acid

This was similarly prepared from Intermediate 37 (0.27mmol) and Intermediate 45 (0.117g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:0.5) to give the title compound as a white solid (0.056g). LCMS: R_t 3.80 min; m/z 765 [$M-H^-$].

Example 50: (2S)-3-(4-[(4-[(2-(4-Chlorophenyl)acetyl)amino]-1-piperidinyl)carbony]oxy}phenyl)-2-((2S)-4-methyl-2-[(2-[(3-(1-piperidinyl)carbonyl)-2-naphthyl]oxy]acetyl)amino]pentanoyl)amino)propanoic acid

This was similarly prepared from Intermediate 37 (0.27mmol) and Intermediate 44 (0.173g). The crude product was purified by flash column chromatography on silica gel eluting with chloroform/methanol/acetic acid (95:5:0.5) to give the title compound as a white solid (0.062g). LCMS: R_t 3.71 min; m/z 868 (MH^+).

Example 51: (2S)-2-[(2S)-2-(2-[2-(Tert-butyl)phenoxy]acetyl)amino]-4-methylpentanoyl]amino}-3-{4-[(4-[(2-phenylacetyl)amino]-1-piperidinyl)carbonyl]oxy}phenyl)propanoic acid

Intermediate 33 (0.23mmol) was treated with 1:1 dichloromethane/THF (3mL). Intermediate 59 (0.105g) was added followed by diisopropylethylamine

(0.16mL). After shaking for 18h at 20°C the resin was filtered, washed with dichloromethane (4 x 5mL) and ether (3 x 5mL) and then dried *in vacuo*. LCMS showed that some of the 4-nitrophenyl carbonate had been hydrolysed to the phenol so the resin was treated with 1:1 dichloromethane/THF (3mL), diisopropylethylamine (0.2mL) and 4-nitrophenyl chloroformate (0.23g). After shaking for 18h at 20°C the resin was filtered and washed with dichloromethane (4 x 5mL) then treated with 1:1 dichloromethane/THF (3mL), Intermediate 59 (0.07g) and diisopropylethylamine (0.12mL). After shaking for 18h at 20°C the resin was filtered and washed with dichloromethane (4 x 5mL) then treated with 1:1 trifluoroacetic acid/dichloromethane (3mL). After 30 mins the resin was filtered and the filtrate was evaporated *in vacuo*. The residue was co-evaporated with dichloromethane followed by ether to give the title compound as an off-white solid (0.083g). LCMS: R_t 3.99 min; *m/z* 729 (MH^+).

15 Example 52: (2S)-2-{[(2S)-2-({2-[2-(Tert-butyl)phenoxy]acetyl}amino)-4-methyl pentanoyl]amino}-3-{4-[(4-[(2-cyclohexylacetyl)amino]-1-piperidinyl]carbonyl}oxy]phenyl}propanoic acid

This was similarly prepared from Intermediate 33 (0.23mmol) and Intermediate 57 (0.106g). The title compound was obtained as an off-white solid (0.073g).
20 LCMS: R_t 4.27 min; *m/z* 735 (MH^+).

Example 53: (2S)-2-{[(2S)-2-({2-[2-(Tert-butyl)phenoxy]acetyl}amino)-4-methyl pentanoyl]amino}-3-{4-[(4-[(2,2-dicyclohexylacetyl)amino]-1-piperidinyl]carbonyl}oxy]phenyl}propanoic acid

25 This was similarly prepared from Intermediate 33 (0.25mmol) and Intermediate 58 (0.144g). The title compound was obtained as an off-white solid (0.105g).
LCMS: R_t 4.63 min; *m/z* 817 (MH^+).

Example 54: (2S)-2-[((2S)-4-Methyl-2-[(2-(2-methylphenoxy)acetyl]amino)pentanoyl)amino]-3-{4-[(4-[(2-phenylacetyl)amino]-1-piperidinyl}carbonyl]oxy]phenyl}propanoic acid

This was similarly prepared from Intermediate 34 (0.3mmol) and Intermediate 59 (0.196g). The crude product was purified by flash column chromatography on silica gel eluting with dichloromethane/methanol/acetic acid/water (240:15:3:2) to give the title compound as a pale yellow foam (0.091g). LCMS: R_t 3.49 min; m/z 687 (MH^+).

10 Example 55: (2S)-2-[((2S)-2-[(2-Cyclohexylphenoxy)acetyl]amino)-4-methylpentanoyl)amino]-3-{4-[(4-[(2-phenylacetyl)amino]-1-piperidinyl}carbonyl]oxy]phenyl}propanoic acid

Intermediate 42 (0.27mmol) was treated with a solution of Intermediate 59 (0.178g) in 1:1 dichloromethane/THF (2mL) followed by diisopropylethylamine (0.95mL). After shaking for 2h at 20°C the resin was filtered and washed with dichloromethane (5 x 5mL) then treated with 1:1 trifluoroacetic acid/dichloromethane (3mL). After 30 mins the resin was filtered and the filtrate was evaporated *in vacuo*. The residue was triturated with ether to give the title compound as an off-white solid (0.074g). LCMS: R_t 4.04 min; m/z 755 (MH^+).

20 Example 56: (2S)-3-{4-[(4-[(2-Cyclohexylacetyl)amino]-1-piperidinyl}carbonyl]oxy]phenyl}-2-[((2S)-2-[(2-(2-cyclohexylphenoxy)acetyl]amino)-4-methylpentanoyl)amino]propanoic acid

This was similarly prepared from Intermediate 42 (0.27mmol) and Intermediate 57 (0.18g). The title compound was obtained as an off-white solid (0.102g). LCMS: R_t 4.22 min; m/z 761 (MH^+).

Biological Data

The compounds of the Examples were tested in assay (1), the Jurkat adhesion assay, and the results obtained were as follows:

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Example	pIC ₅₀	SEM*	n*
1	7.88	0.18	6
2	8.03	0.24	4
3	7.38	0.12	4
4	7.78	0.08	4
5	8.11	0.03	4
6	8.25	0.06	4
7	8.58	0.03	4
8	7.37	0.15	4
9	7.58	0.10	5
10	8.08	0.05	9
11	8.08	0.12	10
12	7.96	0.06	8
13	7.59	0.11	4
14	7.78	0.07	4
15	8.57	0.04	8
16	8.49	0.10	8
17	8.59	0.09	8
18	8.43	0.38	5
19	8.12	0.06	5
20	7.83	0.03	6
21	8.41	0.07	9
22	7.65	0.17	4

Example	pIC ₅₀	SEM*	n*
23	8.35	0.02	10
24	8.22	0.08	10
25	8.50	0.08	10
26	8.53	0.03	4
27	8.55	0.10	7
28	8.46	0.05	10
29	7.79	0.08	6
30	8.24	0.03	4
31	7.59	0.04	4
32	7.62	0.13	6
33	8.46	0.03	9
34	7.57	0.14	4
35	8.18	0.06	6
36	7.91	0.07	6
37	8.24	0.07	6
38	7.81	0.15	4
39	7.65	0.12	4
40	8.04	0.15	4
41	8.03	0.07	4
42	7.96	0.07	6
43	7.65	0.07	6
44	7.62	0.05	5
45	7.24	0.11	6
46	7.36	0.04	4
47	7.48	0.07	4
48	7.38	0.04	4

Example	pIC ₅₀	SEM*	n*
49	7.35	0.06	4
50	7.60	0.10	4
51	7.86	0.05	8
52	7.48	0.21	4
53	6.81	0.10	5
54	8.25	0.03	5
55	7.21	0.13	4
56	7.06	0.19	6

*SEM standard error of the mean of n experiments

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Abbreviations

WSCDI	1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
PyBop	benzotriazol-1-yl-oxy-trispyrrolidinophosphonium hexafluorophosphate
DIC	1,3-diisopropylcarbodiimide
HOBT	1-hydroxybenzotriazole
Boc	tert butoxycarbonyl
Fmoc	9-fluorenylmethoxycarbonyl
Cbz	carbobenzyloxy
DIPEA	diisopropylethylamine
DCM	dichloromethane
DMF	dimethylformamide
THF	tetrahydrofuran

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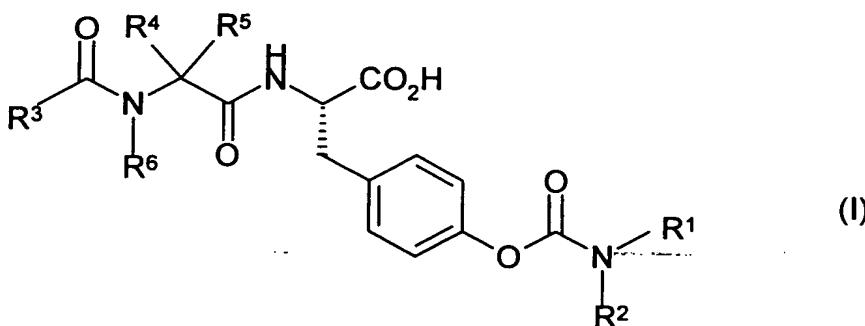
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The application of which this description and claims forms part may be used as a basis for priority in respect of any subsequent application. The claims of such subsequent application may be directed to any feature or combination of features described herein. They may take the form of product, composition, process or use claims and may include, by way of example and without limitation, one or more of the following claims.

CLAIMS

1. A compound of formula I:



5

wherein R¹ and R² independently represent

(i) -C₁₋₆ alkyl, -C₃₋₈ cycloalkyl or -C₁₋₃ alkylC₃₋₈ cycloalkyl,

or such a group in which alkyl or cycloalkyl is substituted by one or more

10 halogen, -CN, nitro, hydroxy or -OC₁₋₆alkyl groups;

(ii) -(CH₂)_eAr¹ or -(CH₂)_eOAr¹;

or NR¹R² together represent pyrrolidinyl, piperidinyl, piperazinyl, thiomorpholinyl, morpholinyl or azepinyl, or such a group fused to a benzene ring, optionally substituted by one or more -(CO)_n(CH₂)_tAr¹, -(CO)_nC₁₋₆ alkylAr¹Ar²,

15 -(CO)_nC₁₋₆alkyl, -(CH₂)_rOH, -(CH₂)_rO(CH₂)_pOH, -(CH₂)_rOC₁₋₆ alkyl, -O(CH₂)_tAr¹,

-(CH₂)_rSO₂Ar¹, piperidin-1-yl, -(CH₂)_tCONR⁸R⁹, -NR¹⁰(CO)_n(CH₂)_tAr¹,

-NR¹⁰(CO)_nC₁₋₃alkylC₃₋₆ cycloalkyl, -NR¹⁰(CO)_nC₁₋₆ alkylidic₃₋₆ cycloalkyl,

-CONR¹⁰(CH₂)_tAr¹, halogen, -NHSO₂C₁₋₆alkyl, -SO₂NR¹⁰R¹¹, -SO₂C₁₋₆ alkyl or

-SO₂Ar² groups;

20

R³ represents -C₁₋₆alkylNHC(=NH)NH₂, -C₂₋₆alkenylNHC(=NH)NH₂,

-C₂₋₆alkynylNHC(=NH)NH₂, -C₁₋₆alkylNR¹⁴R¹⁸, -(CH₂)_hCONR¹⁴R¹⁸,

-(CH₂)_hCOC₁₋₆alkyl, -(CH₂)_dCHNR¹⁸CONR²⁰R²¹, -(CH₂)_mNR¹⁸CONR¹⁴R¹⁸,

$-(CH_2)_dNR^{18}Ar^3$, $-(CH_2)_dCONR^{18}Ar^3$, $-(CH_2)_hCOOR^{18}$, $-(CH_2)_cAr^3$, $-O(CH_2)_cAr^3$,

$-(CH_2)_dCO(CH_2)_sAr^3$ or $-(CH_2)_dOAr^3$;

or R^3 represents $-(CH_2)_c$ -2,4-imidazolidinedione, $-(CH_2)_c$ (piperidin-4-yl),

$-(CH_2)_c$ (piperidin-3-yl), $-(CH_2)_c$ (piperidin-2-yl), $-(CH_2)_c$ (morpholin-3-yl) or

5 $-(CH_2)_c$ (morpholin-2-yl) optionally substituted on nitrogen by $-(CO)_fC_{1-6}$ alkyl,

$-(CO)_f(CH_2)_cAr^2$ or $-C(=NH)NH_2$;

or R^3 represents $-(CH_2)_z$ dibenzofuran optionally substituted by $-C_{1-6}$ alkyl or

halogen;

or R^3 represents $-(CH_2)_c$ -thioxanthen-9-one;

10

R^4 represents hydrogen, $-C_{1-6}$ alkyl, $-C_{1-3}$ alkyl C_{3-6} cycloalkyl, $-(CH_2)_qAr^2$,

$-C_{1-4}$ alkyl-X-R⁷, $-C_{1-4}$ alkyl SO₂C₁₋₄ alkyl, $-C_{1-6}$ alkylNR¹²R¹³ or

$-C_{1-6}$ alkylNR¹²COC₁₋₆ alkyl;

15

R^5 represents hydrogen, or R^4R^5 together with the carbon to which they are attached form a C₅₋₇ cycloalkyl ring;

R^6 represents hydrogen or $-C_{1-6}$ alkyl, or R^6 and R^4 together with the N and C atoms to which they are respectively attached form a pyrrolidine ring;

20

R^7 represents hydrogen, $-(CH_2)_wNR^{12}R^{13}$, $-(CH_2)_uAr^2$ or

$-(CH_2)_wNR^{12}COC_{1-6}$ alkyl;

R^8 , R^9 , R^{16} and R^{17} independently represent hydrogen, $-C_{1-6}$ alkyl, $-C_{3-6}$ cycloalkyl,

25 $-C_{1-3}$ alkyl C_{3-6} cycloalkyl, $-C_{2-6}$ alkenyl or NR⁸R⁹ or NR¹⁶R¹⁷ together represents morpholinyl, pyrrolidinyl, piperidinyl, piperazinyl or piperazinyl N-substituted by $-C_{1-6}$ alkyl, -COphenyl or -SO₂methyl;

R^{10} , R^{11} , R^{12} , R^{13} , R^{15} , R^{18} , R^{20} and R^{21} independently represent hydrogen or $-C_{1-6}\text{alkyl}$;

5 R^{14} , R^{19} and R^{22} independently represent hydrogen, $-C_{1-6}\text{alkyl}$, $-C_{3-6}$ cycloalkyl or $-(CH_2)_x Ar^4$ or $NR^{14}R^{18}$ or $NR^{15}R^{22}$ together represents morpholinyl, pyrrolidinyl, piperidinyl, piperazinyl or $N-C_{1-6}\text{alkylpiperazinyl}$;

Ar^1 represents phenyl or a 5 or 6 membered heterocyclic aromatic ring containing 1 to 3 heteroatoms selected from O, N and S optionally substituted by one or more halogen, $C_{1-6}\text{alkyl}$, hydroxy, $-OC_{1-6}\text{alkyl}$, CF_3 , nitro, $-Ar^2$ or $-OAr^2$ groups;

Ar^2 represents phenyl optionally substituted by one or more halogen, $-C_{1-6}\text{alkyl}$, hydroxy, $-OC_{1-6}\text{alkyl}$, $-CF_3$ or nitro groups;

15

Ar^3 represents phenyl, a 5 or 6 membered heterocyclic aromatic ring containing 1 to 3 heteroatoms selected from O, N or S, or such a group fused to a benzene ring, optionally substituted by one or more $-CO(CH_2)_g Ar^4$, $-(CH_2)_y Ar^4$, $-(CH_2)_y COAr^4$, $-(CO)_a C_{1-6}$ alkyl, $-(CO)_a C_{2-6}$ alkenyl, $-(CO)_a C_{2-6}$ alkynyl,

20 $-(CO)_a C_{3-6}$ cycloalkyl, $-(CO)_a C_{1-6}$ haloalkyl, halogen, $-COCH_2 CN$, $-(CH_2)_b NR^{16}R^{17}$, $-(CH_2)_b NHC(=NH)NH_2$, $-CYNR^{16}(CO)_a R^{17}$, $-(CH_2)_b NR^{15} COR^{19}$, $-(CH_2)_b CONR^{15}R^{22}$, $-(CH_2)_b NR^{15} CONR^{15}R^{22}$, $-(CH_2)_b CONR^{15}(CH_2)_j NR^{15} R^{22}$, $-(CH_2)_b SO_2 NR^{15} R^{22}$, $-(CH_2)_b SO_2 NR^{15} COAr^2$, $-(CH_2)_b NR^{15} SO_2 R^{19}$, $-SO_2 R^{19}$, $-SOR^{19}$, $-(CH_2)_z OH$, $-COOR^{15}$, $-CHO$, $-OC_{1-10}\text{alkyl}$, $-O(CH_2)_j NR^{15} R^{22}$,

25 $-O(CH_2)_j NHC(=NH)NH_2$, $-O(CH_2)_b CONR^{16}R^{17}$, $-O(CH_2)_k COOR^{15}$, $-O(CH_2)_j OAr^2$, $-O(CH_2)_b Ar^2$, 3-phenyl-2-pyrazolin-5-one or 4,5-dihydro-3(2H)-pyridazinone groups;

Ar⁴ represents phenyl or a 5 or 6 membered heterocyclic aromatic ring containing 1 to 3 heteroatoms selected from O, N and S optionally substituted by one or more halogen, -C₁₋₆alkyl, hydroxy, -OC₁₋₆alkyl, -CF₃, nitro or -CONH₂ groups;

5

X and Y independently represent O or S;

a, f, k, s and n independently represent 0 or 1;

b, c, r, x, y and z independently represent an integer 0 to 2;

10 d, g and u independently represent 1 or 2;

e, h, q and w independently represent an integer 1 to 3;

j and p independently represent an integer 2 to 4;

m independently represents an integer 0 to 4;

t independently represents an integer 0 to 3;

15 and salts and solvates thereof.

2. A pharmaceutical composition comprising a compound of formula (I) as defined in claim 1 or a pharmaceutically acceptable salt or solvate thereof in admixture with one or more pharmaceutically acceptable diluents or carriers.

20

3. A compound of formula (I) as defined in claim 1 or a pharmaceutically acceptable salt or solvate thereof for use as a pharmaceutical.

25 4. Use of a compound of formula (I) as defined in claim 1 or a pharmaceutically acceptable salt or solvate thereof in the manufacture of a medicament for the treatment of inflammatory diseases.

5. A method of treatment or prophylaxis of inflammatory diseases eg. asthma which comprises administering to a patient an effective amount of a

compound of formula (I) as defined in claim 1 or a pharmaceutically acceptable salt or solvate thereof.

